

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw mode:

Run on: October 24, 2003, 15:43:05 ; Search time 2765.41 seconds
(without alignments): 14602.112 Million cell updates/sec

Title: US-09-830-902-1_COPY_100000_101000

Perfect score: 1001

Sequence: 1 aatataaaatgcataacttt.....acctgattctgtctctctttt 1001

Scoring table: IDENTITY NJC

Gapop 10.0 , Gapext 1.0

Searched: 2898711 seqs, 2045481386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:

1: gb_ba.*

2: gb_hg.*

3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pa.*

7: gb_ph.*

8: gb_pi.*

9: gb_pr.*

10: gb_ro.*

11: gb_sts.*

12: gb_sy.*

13: gb_un.*

14: gb_va.*

15: em_ba.*

16: em_fun.*

17: em_hum.*

18: em_in.*

19: em_mu.*

20: em_om.*

21: em_or.*

22: em_ov.*

23: em_pat.*

24: em_ph.*

25: em_pa.*

26: em_ro.*

27: em_sts.*

28: em_un.*

29: em_va.*

30: em_hg_hum.*

31: em_hg_inv.*

32: em_hg_other.*

33: em_hg_mus.*

34: em_hg_pin.*

35: em_hg_rtd.*

36: em_hg_mam.*

37: em_hg_vit.*

38: em_sy.*

39: em_higo_hum.*

40: em_higo_mus.*

41: em_higo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than, or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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3	1001	100.0	162692	9	CNS01DS8	AL21659 BAC sequ
4	1001	100.0	185281	2	AC011232	AC011232 Homo sapi
5	403	40.3	3263	6	AX093472	AX093472 Sequence
6	403	40.3	3263	9	HS24460C1	AC02460C1 Homo sapi
7	403	40.3	5120	9	ABC29006	ABC29006 Homo sapi
8	200.6	20.0	172816	9	ABC293899	ABC293899 Homo sapi
9	199.8	20.0	195432	9	AC130403	AC130403 Homo sapi
10	199.8	20.0	203354	2	ACC68752	ACC68752 Homo sapi
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32	192.8	19.3	140363	9	AC142331	AC142331 Par trogl
33	192.8	19.3	172533	9	AC002381	AC002381 Human BAC
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ALIGNMENTS

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LOCUS AX093471 110000 bp DNA linear PAT 30-MAR 2001
DEFINITION Sequence 1 from Patent WO0118198.
ACCESSION AX093471
VERSION AX093471.1 GI:13509911
KEYWORDS Homo sapiens (human)
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
REFERENCE
AUTHORS Weissbach, J. and Hazan, J.
TITLE Cloning, expression and characterization of the spg4 gene
responsible for the most frequent form of autosomal spastic


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TITLE      Spectrum of SP4 mutations in autosomal dominant spastic paraplegia
JOURNAL    Hum. Mol. Genet. 9 (4), 637-644 (2000);
MEDLINE    20164302
PUBMED     10699187
REFERENCE  3 (bases 1 to 110000);
AUTHORS    Direct Subrission
TITLE      Submitted (17-JUN-1999) Genoscope, Genoscope - Centre National de
JOURNAL    Sequencing, BP 197, EVRY 91056, FRANCE
COMMENT    E-mail : seqref@genoscope.cns.fr - Web : www.genoscope.cns.fr. The
          sequence is the result of the assembly of 2 BAC clones: R-336p14
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JOURNAL      Nat. Genet. 23 (3), 296-303 (1999)
MEDLINE      20055425
PubMed      10610178
REFERENCE    2
AUTHORS      Focke, N., Navel, J., Byrne, P., Davoine, C., Crnaud, C.,
              Boentsch, D., Samson, C., Coulinho, P., Hutchinson, M., McMonagle, P.,
              Burgunder, C., Tarfiglione, A., Heitzler, O., Fekli, I., Doufel, T.,
              Parfrey, N., Brice, A., Fontaine, B., Prud'homme, C., Weissenbach, J.,
              Durr, A. and Hazan, J.
TITLE        Spectrum of SPG4 mutations in autosomal dominant spastic paraplegia
JOURNAL      Hum. Mol. Genet. 9 (4), 637-644 (2000)
MEDLINE      20164322
PubMed      10699187
REFERENCE    3 (bases 1 to 3263)
AUTHORS      Genoscope.
JOURNAL      Submitted (17-JUN-1999) Genoscope, Genoscope - Centre National de
              Sequencage, BP 191, EVRY 91006, FRANCE
COMMENT      E-mail : secre@genoscope.cns.fr - Web : www.genoscope.cns.fr.
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DB 1852 AGATGAGAAATATTCGATTATCTGACTTCTACTGAACTTCCTTGAAGAAATATAAAGCGAGCG 1911
QY 659 TCAGCCCTCAAACTTTAGAGCGGTACATCGTTTGAACGAGGACTTTGAGATACCACTG 718
DB 1912 TCAGCCCTCAAACTTTAGAGCGGTACATCGTTTGAACGAGGACTTTGAGATACCACTG 1971
QY 719 TTTAAGGAAATACCTTTGTAAACCTCGAGAACATTTTACTTAAAGAGGAAACACAGAT 778
DB 1972 TTTAAGGAAATACCTTTGTAAACCTCGAGAACATTTTACTTAAAGAGGAAACACAGAT 2031
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QY 839 ACATATTGTGCCACCAACTTGAAGATGAACCCAGAACACAGACTTTAAACAAATATACAAA 898
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QY 959 AACTACTAGTTAGAGCACACAAACCTGATTTGCTGCTTCTCTT 1001
DB 2212 ACATAAGTTAGAGCACACAAACCTGATTTGCTGCTTCTCTT 2254

RESULT 7
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LOCUS      AB029006 Homo sapiens mRNA for KIAA1083 protein, complete cds.
DEFINITION
ACCESSION AB029006
VERSION    AB029006.1 GI:5689502
KEYWORDS
SOURCE     Homo sapiens (human)
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            1 (sites)
REFERENCE 1 (bases 1 to 5120)
AUTHORS    Kikuno, R., Nagase, T., Ishikawa, K., Hirose, M., Miyajima, N.,
            Tanaka, A., Kotani, H., Nomura, N. and Ohara, O.
TITLE      Prediction of the coding sequences of unidentified human genes.
            XIV. The complete sequences of 100 new cDNA clones from brain which
            code for large proteins in vitro
JOURNAL    DNA Res. 6 (3), 197-205 (1999)
MEDLINE    99397452
PubMed     10478851
REFERENCE 2 (bases 1 to 5120)
AUTHORS    Chara, O., Nagase, T. and Kikuno, R.
TITLE      Direct Submission
JOURNAL    Submitted (17-JUN-1999) Osamu Chara, Kazusa DNA Research Institute,
            Laboratory of DNA Technology, Yana 1512-3, Kisarazu, Chiba
            232-0812, Japan (E-mail:cdnaif@kazusa.or.jp, Tel:+81-438-52-3913,
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Query Match 40.3%; Score 403; DB 9; Length 5120;
Best Local Similarity 100.0%; Pred. No. 5,5e-67;
Matches 403; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 599 AGATGAGAAATATTCGATTATCTGACTTCTACTGAACTTCCTTGAAGAAATATAAAGCGAGCG 658
DB 1852 AGATGAGAAATATTCGATTATCTGACTTCTACTGAACTTCCTTGAAGAAATATAAAGCGAGCG 1911
QY 659 TCAGCCCTCAAACTTTAGAGCGGTACATCGTTTGAACGAGGACTTTGAGATACCACTG 718
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QY 719 TTTAAGGAAATACCTTTGTAAACCTCGAGAACATTTTACTTAAAGAGGAAACACAGAT 778
DB 1972 TTTAAGGAAATACCTTTGTAAACCTCGAGAACATTTTACTTAAAGAGGAAACACAGAT 2031
QY 779 CTTCATGACGCTCATCGGCTAGAGAACACGCTTACGAGCTTTTAGAGTCTT 839
DB 2032 CTTCATGACGCTCATCGGCTAGAGAACACGCTTACGAGCTTTTAGAGTCTT 2091
QY 839 ACATATTGTGCCACCAACTTGAAGATGAACCCAGAACACAGACTTTAAACAAATATACAAA 898
DB 2092 ACATATTGTGCCACCAACTTGAAGATGAACCCAGAACACAGACTTTAAACAAATATACAAA 2151

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QY 959 ACATAGATTAGACACAAAGCAACACCTGATCTGGCTTCCTT 1001
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RESULT 8
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LOCUS AC093899
DEFINITION Homo sapiens BAC clone RP11-724016 from 2, complete sequence.
ACCESSION AC093899
VERSION AC093899.1 GI:18497265
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SOURCE Homo sapiens (human);
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 172816)
AUTHORS Suiston,J.E. and Waterston,R.
TITLE Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1108 (1998).
MEDLINE 98063782
PubMed 9847074
REFERENCE 2 (bases 1 to 172816)
AUTHORS Pearman,C., Haakensen,W. and Boyer,E.
TITLE The sequence of Homo sapiens BAC clone RP11-724016
JOURNAL Unpublished (2001)
REFERENCE 3 (bases 1 to 172816)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (10-SEP-2003) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE 4 (bases 1 to 172816)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (05-FEB-2002) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE 5 (bases 1 to 172816)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (12-JUN-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Feb 5, 2002 this sequence version replaced gi:15625013.
COMMENT ----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: saplens@wustl.wustl.edu
----- Summary Statistics
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Center project name: H_NH0724016
Drafting Center: WIBR

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping core sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality > 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RP11 human BAC library was made from the blood of one male donor, as described by Osoegawa,K., Woon,P.Y., Zhao,B., Frengen,E., Ratens,M., Catanese,J.C. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genetics 51:1-9. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at <http://www.chori.org>

VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:

Actual start of this clone is at base position 1 of RP11-724016; actual end is at base position 172816 of RP11-724016.

Unresolved simple sequence repeat from base position 22700 to 25900.

The sequence of AC093894 has been incorporated into AC093899.

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organism	"Homo sapiens"
molecule	"genomic DNA"
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Query Match 20.0%; Score 200.6; DB 9; Length 172616;
Best Local Similarity 60.8%; Pred. No. 1,2e-28;
Matches 398; Conservative 0; Mismatches 249; Indels 9; Gaps 4.

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Db 24223 TTATAATTATCATTTTGGCTGGGCGCACTGGCTCATCTCTGTAAATCCGAGCATTGGG 26164
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Db 26163 AGCTGAGTGGGCGGATCATCTGAGTTCAGGAGTTCGAGACCAACCTGGCCATCATG 26104
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RESULT 9
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DEFINITION Human BAC Library) complete sequence.
ACCESSION AC130403
VERSION AC130403.3 GI:24431574
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SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

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REFERENCE
AUTHORS

- 1 (bases 1 to 195432)
 Vuzny D.M., Adams C., Adio-Oduola B., Ali-Saman F.R., Allen C., Ambrosio S.L., Amaral-Engel H.C., Are, C.R., Ayala M., Banks T., Barbara J., Benton C., Benge K., Blankenburg K., Bonnin D., Buck J., Bowie S., Brieva M., Brown E., Brown M., Bryant N.P., Burch C., Burch P., Burkett C., Burrell K.L., Byrd N.C., Cayton T.F., Carter M., Cavazos S.R., Chacko J., Chavez D., Chen G., Chen R., Chen Z., Chiu D., Chowdhry I., Christopoulos C., Cleveland C.D., Cox C., Coyle M.D., Dathore S.B., David R., Davila M.L., Davis C., Davy-Carroll L., Federich P.A., Deane J., Davis R., Deigado J., Denn A.L., Ding Y., Dinch H.R., Douthwaite K.J., Draper H., Dugan-Rocha S., Durkin K.J., Earnhart C., Edgar D., Edwards C.C., Elhai C., Ewelling J., Escoto C., Falls T., Ferraguto D., Flagg N., Ford J., Foster P., Frantz P., Gabis A., Gao J., Garcia A., Garner T., Garza N., Gill R., Gorelli C.H., Guevara W., Gunaratne P., Hale S., Hamilton K., Han J., Harris K., Harris K., Hart M., Havlak P., Hayes A., Hernandez J., Hernandez C., Hodgson A., Hogue M., Holoway C., Hollins E., Homai P., Howard S., Hubert S., Hulyk S., Hu-e J., Ioshikhes I., Jackson J.E., Jacobson B., Jia Y., Johnson R., Jolivet S., Joudah S., Karlsson E., Kelly S., Khan S., King L., Koryah J., Kovar C., Kratochvil J., Kureshi A., Landry N., Leal B., Lee E., Lewis L.C., Lewis S., Liang J.Z., Lucharske D., Liu C., Liu J., Liu N., Louiscad R., Lora D., Lora J., Lora X., Lucier A., Lucier R., Luna R., Ma J., Maheshwari N., Massey E., Marcandell I., Martin R., Martindale A., Martinez E., Massey E., Mathiney E., McLeod M.P., Meador M., Mei G., Merscher S., Metzger M., Miller A., Miner G., Miner Z., Mitchell T., Mohabhat K., Montgomery K.T., Moragan M., Morris S., Moser V., Neal D., Neilson D., Newton J., Newton N., Nguyen A., Nguyen N., Nguyen N., Nickerson E., Nwkekwu S., Ogburn C., Okwuonu S., Oreguene N., Oviedo R., Pace A., Payton B., Peery J., Perez J., Peters L., Pickens R., Primus E., Pu L.L., Quiles M., Ren Y., Rivas M., Rojas A., Rojebakar L., Roife M., Ruiz S., Savory G., Scherer S., Scott G., Shen H., Shim C., Shostakov N., Sisson I., Sodergren E., Soroka T., Sparks A., Stanley H., Stone H., Sutton A., Svatek A., Tabor P., Tamerisa A., Tamerisa K., Tang R., Tansey J., Taylor C., Taylor T., Teifrod P., Thomas N., Thomas R., Ustani K., Vasquez J., Vera V., Villalón D., Vinson R., Wang Q., Wang S., Ward-Moore S., Warren R., Washington C., Wallington S., Williams G., Williamson A., Wleczek R., Wooden S., Worley K., Wu C., Wu Y., Wu Y.F., Zhou J., Zorrilla S., Kuchelapati R., Weinstein G. and Gibbs R.
- Direct Submission
 Unpublished
 2 (bases 1 to 195432)
 Worley K.C.
- Direct Submission
 Submitted (10-AUG-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
- 3 (bases 1 to 195432)
 Worley K.C.
- Direct Submission
 Submitted (11-OCT-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
- 4 (bases 1 to 195432)
 Worley K.C.
- Direct Submission
 Submitted (31-OCT-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
- 5 (bases 1 to 195432)
 Worley K.C.
- Direct Submission
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- On Oct 31, 2002 this sequence version replaced GI23803880.
 INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email
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FEATURES

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RESULT 11
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DEFINITION
complete sequence.
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VERSION AC002549.1 G1:2992475
KEYWORDS HTO.
SOURCE   Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 163027)
Murny,D., Aronson,A.D., Brundage,E., Carvelli,K., Chen,E., Chen,C.,
Di,W., Ding,Y., Dugan,S., Durbin,J., Forcum,J., Ganesh,R.,
Garcia,C., Goodman,M., Gorrell,J.H., Haywood,W., Jackson,J.,
Jin,S., Karpal,R., Karpathy,S., Leal,B., Li,Y., Liu,W., Logan,C.,
Lu,C., Ly,T., Martinez,C., Oswal,G., Perez,J., Rashid,N.D., Logan,C.,
Rowland,K., Savage,L., Scherer,S., Shen,H., Timms,X.M., Todd,C.,
Vo,O., Worley,K.C., Yu,W., Chinault,C., Nelson,D. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 163027)
Chiu,X.W.
Direct Submission
Submitted (18-SEP-1997) Molecular and Human Genetics, Baylor
College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 163027)
Worley,K.C.
Direct Submission
Submitted (27-MAR-1998) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Mar 27, 1998 this sequence version replaced g1:2960313.
Sequencing is completed to a minimum standard of double strand
coverage with a minimum of 2 clones and 2 reads with no ambiguities
or 2 chemistries with a minimum of 2 clones and 3 reads with no
ambiguities. If the sequence quality does not meet this standard,
it will be indicated in the annotation.

The repeat regions shown were identified using RepeatMasker by
Adrian S-It.

Sequence similarities were identified using PowerBlast by Jinghui
Zhang.

Exon/intron boundaries of identified genes were chosen if there
were canonical splice junctions that maintained sequence continuity
across the splice junctions.
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REFERENCE
AUTHORS

3 (bases 1 to 119922)

Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Bouckghalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., DeArrelano,K., Dewar,K., Diaz,J.S., Dodge,S.,
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Karakas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
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Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schupback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talaras,J.,
Tesfaye,S., Theodore,J., Topham,K., Travers,Y., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,X.

TITLE
JOURNAL

Submitted (21-AUG-2002): Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA

4 (bases 1 to 119922)

REFERENCE
AUTHORS

Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Bouckghalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
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TITLE
JOURNAL

Submitted (28-AUG-2002): Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA

COMMENT

On Aug 28, 2002 this sequence version replaced gi:22391567.

All repeats were identified using RepeatMasker.

Smith, A.F.A. & Green, P. (1996-1997)

<http://ftp.genome.washington.edu/RX/RepeatMasker.html>

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seg.wi.mit.edu>

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: 110689

Center clone name: 2357_A8

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RESULT 15
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LOCUS
DEFINITION Human DNA sequence from clone RP4-738A13 on chromosome
X21:31-22.1, complete sequence.
ACCESSION AL109801
VERSION AL109801.14 GI:281933390
KEYWORDS H.G.
SOURCE Homo sapiens
ORGANISM Homo sapiens (human);
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Whitehead, S.
Direct Submission
Submitted (31-JAN-2003) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail: enquiries@
humquerry@sanger.ac.uk
On Feb 1, 2003 this sequence version replaced GI:6624916.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
EMBL, EMBL; SWISSPROT, TrEMBL; TrEMBL; WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome X, constructed by the Sanger Centre Chromosome X Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/chrX
----- Genome Center
Center: Wellcome Trust Sanger Institute

```

Center code: SC
Web site: <http://www.sanger.ac.uk>
Contact: humquerry@sanger.ac.uk

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC.
RP4-738A13 is from the library RPCr-4 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>
VECTOR: pCYPAC2.

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Db 39666 AACCTGTCTCTACTAAAAATACAAAAATAGCTGGGATGGAGGATGTGCTGTAAATC 39607
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 187 CCAGCTACTCGGAGGCTGAGACAGAGAAATGCTTGAACCTTGAGGGGAGGATTGCA 245
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 39606 CCAGCTACTCGGAGGCTGAGACAGAGAAATGCTTGAACCTTGAGGGGAGGATTGCA 39548
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Search completed: October 24, 2003, 21:06:17
Job time : 2770.61 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw mode:

Run on: October 24, 2003, 15:39:20 : Search time 252.249 seconds
(without alignments)
10722.169 Million cell updates/sec

Title: US-09-830-902-1_COPY_100000_101000

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Sequence: 1 aatataaaatgcatactctt.....acctgattctggtctctttt tctt

Scoring table: ICENTITY NUC

Gapp 10.0 , Gapext 1.0

Searched: 2552756 seqs, 134979017 residues

Total number of hits satisfying chosen parameters: 5165512

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N Geneseq 19Jun03:*

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- 25: /SIDSL/gcgdata/geneseq/geneseq-emb1/NA2004.DAT:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	100.0	100.0	10000	22	AA584800
2	403	40.3	3263	22	AA584801
3	403	40.3	4152	24	AB054772
4	191.4	19.1	3143	24	AA517746
5	191.4	19.1	3573	22	AA104148
6	191.4	19.1	32195	22	AA104180
7	90.8	19.1	7677	23	AB23205
8	190.8	19.1	7677	23	AB232052

C	9	190.4	19.0	3074	21	AA576994	Human CREX ORF2549
C	10	189.8	19.0	10360	18	AA576164	Human alpha-N-acet
C	11	189.8	18.9	7792	22	AA532727	Human genomic DNA
C	12	189.8	18.9	7792	22	AA535099	Human cardiovascular
C	13	189.6	18.9	19516	22	AA150294	Human nervous syst
C	14	189.6	18.9	19516	22	AA158556	Human nervous syst
C	15	189.6	18.8	19516	22	AA136855	Human musculoskele
C	16	189.6	18.8	19516	25	ABX59843	cDNA encoding nove
C	17	187.9	18.8	50200	21	AA596166	Poly-crinic repeat
C	18	187.6	18.7	4768	21	AA569134	Human ABC1 gene ex
C	19	187.6	18.7	11754	22	AA504033	Human ABC1 gene, p
C	20	187.6	18.7	11754	24	ABJ58149	Human ABCA1 transp
C	21	187.6	18.7	52616	22	AAK70459	Human immune/haema
C	22	187.6	18.7	52616	22	AAK78930	Human immune/haema
C	23	187.6	18.7	97835	24	AEK84796	Human CDNA difere
C	24	187.6	18.7	183959	22	AAF92831	Human ABC1 genomic
C	25	187	18.7	3417	22	AAH18467	Human CDNA sequenc
C	26	186.6	18.6	512	21	AA574416	Human secreted pro
C	27	186.6	18.6	6216	22	AAK81014	Human immune/haema
C	28	186.6	18.6	9469	22	AAK79514	Human immune/haema
C	29	186.6	18.6	17335	23	ABK42193	Genomic sequence #
C	30	186.6	18.6	19882	23	ABK42394	Genomic sequence #
C	31	186.6	18.6	79528	24	AA150614	Human cancer statu
C	32	186.6	18.6	116840	25	ABQ72451	Human VWF DNA, HO
C	33	186.6	18.6	169380	24	ABX58336	Human phosphodiester
C	34	186.4	18.6	2233	22	AAK77256	Human immune/haema
C	35	186.4	18.6	13467	22	AA105944	Human reproductive
C	36	186.4	18.6	13467	22	AA527670	DNA encoding novel
C	37	186.4	18.6	13467	22	AA527838	DNA encoding novel
C	38	186.4	18.6	13467	22	AAK85539	Human immune/haema
C	39	186.4	18.6	13467	21	AB198508	Human testicular a
C	40	186.2	18.6	14491	25	AB209914	Human 5' and/or re
C	41	185.9	18.6	32190	22	AA536709	Human cardiovascular
C	42	185.6	18.5	13058	22	AA529194	Genomic sequence #
C	43	185.6	18.5	13058	22	AAK79125	Human immune/haema
C	44	185.6	18.5	13058	24	AB568334	Human DNA-binding
C	45	185.6	18.5	62804	24	ABD39317	Human calcium/calr

ALIGNMENTS

RESULT 1	
AA584800	
ID	AA584800 standard; DNA; 110000 BP.
XX	AA584800:
AC	AA584800:
XX	09-JUL-2001 (first entry):
XX	Nucleotide sequence of the human SPG4 gene.
XX	Human: SPG4 gene; spactin; PSF-AD; gene therapy;
XX	autosomal dominant familial spastic paraplegia; ss.
XX	Hom sapiens.
XX	Key
XX	Location/Qualifiers
FT	9932..102009
FT	/tag= a
FT	/note= "contains introns"
FT	9932..10471
FT	/tag= b
FT	/number= "1"
FT	10472..33718
FT	/tag= c
FT	/number= "1"
FT	33719..33805
FT	/tag= d
FT	/number= "2"
FT	33806..35748
FT	/tag= e
FT	/number= "2"

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PT /number= "15"
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PT /*tag= ae
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PT 93444..93484
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PT 93485..100599
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PT /*tag= "16"
PT 100600..102009
PT /*tag= ah
PT /*tag= "17"
XX
XX FR2798138-A1.
XX
XX 09-MAR-2001.
XX
XX 03-SEP-1999; 99FR-0011097.
XX
XX 03-SEP-1999; 99FR-0011097.
XX
XX (CNRS : CNRS CENT NAT RECH SCI.
XX
XX Weissenbach J, Hazan J;
XX
XX WPI; 2001-283966/30.
XX
XX New human nucleic acid from the SPG4 gene, useful e.g. for diagnosis of
XX autosomal dominant familial spastic paraplegia and in drug screening .
XX
XX Claim 2: Page 45-106: 145pp; French.
XX
XX The present sequence represents a human SPG4 gene. The SPG4 gene encodes
XX a spastin polypeptide. Mutations in the SPG4 gene are responsible for
XX autosomal dominant familial spastic paraplegia. SPG4 polynucleotides,
XX and their fragments, are used to screen DNA banks for sequences that
XX encode spastin. Particularly sequences in other mammals, specifically
XX mice, to identify SPG4 mutations, or other genetic anomalies,
XX particularly for diagnosis of autosomal dominant familial spastic
XX paraplegia (PSP-AD); to identify promoters and other regulatory elements
XX of the SPG4 gene; for detection and amplification; for recombinant
XX production of spastin; and for diagnostic genotyping of PSP-AD.
XX
XX Sequence 110000 BP; 30622 A; 21640 C; 22817 G; 34921 T; 0 other:
XX
XX Query Match :00.0%; Score 1001; DB 22; Length 110000;
XX Best Local Similarity :00.0%; Pred. No. 2e-171;
XX Matches 1001; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Qy 1 AATATAAAATGCATCTTAGCTGGGCAAGTGGCTCACCTCTGTATATCCAGCACTT 40
D5 |||||
D5 100000 AATATAAAATGCATCTTAGCTGGGCAAGTGGCTCACCTCTGTATATCCAGCACTT 100059
Qy 61 TGGAGGCCCAAGCTGGGCGGATCACCTGAGGTAGAGTTCAGGACCACTGGGCATCA 120
D5 |||||
D5 100060 TGGAGGCCCAAGCTGGGCGGATCACCTGAGGTAGAGTTCAGGACCACTGGGCATCA 100119
Qy 121 TGGGCAAAACCTCTCTCTACTTAAATAACAAAAATTAGCTGGGCATGGAGGCGATGGCT 180
D5 |||||
D5 100120 TGGGCAAAACCTCTCTCTACTTAAATAACAAAAATTAGCTGGGCATGGAGGCGATGGCT 100179
Qy 181 GTATCCAGCTACTCGGGAGCTGAGACAGAGAGTTCCTTGAACCTGGAGGGGGAGG 240
D5 |||||
D5 100180 GTATCCAGCTACTCGGGAGCTGAGACAGAGAGTTCCTTGAACCTGGAGGGGGAGG 100239
Qy 241 TTGCATATCTGAGTGGTGAATTTGTGATCTCTTTTCTCTTTTGTCTGTATTTTGAAC 300
D5 |||||
D5 100240 TTGCATATCTGAGTGGTGAATTTGTGATCTCTTTTCTCTTTTGTCTGTATTTTGAAC 100299
Qy 301 TTCTATAAATGATCTGTGTTTGTGTTTATATATGGAAAAATATATGGTTTCAATGTTA 360
D5 |||||

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DB 100300 TTTCTAATGATTTGTTTCTTTTATATTGATTTGAAATATATATGCTTCTTAAATGTTA 100359
QY 361 ATAGCTATGAACCTAAACACAGTAATATATATATAGCTATAGCTATTTATAGGTTTCT 420
DB 100360 ATACTATGAACCTAAACACAGTAATATATATATATATATATATATATATATATATAGGTTTCT 100419
QY 421 TGTGTAGCAGATACATATAGAAATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 480
DB 100420 TGTGTAGCAGATACATATAGAAATATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 100479
QY 481 ATACAGGTATATTTTATATACATTTAGAAACACAGCAGCATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 540
DB 100482 ATACAGGTATATTTTATATACATTTAGAAACACAGCAGCATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 100539
QY 541 TCGTTAAACCAATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 600
DB 100540 TCGTTAAACCAATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 100599
QY 601 ATGAGATATATCGATTTATCTGACTTCACTGAATCTGTGAAATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 660
DB 100600 ATGAGATATATCGATTTATCTGACTTCACTGAATCTGTGAAATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 100659
QY 661 AGCCCTCAAACTTTAGAGCGTACATATAGCTTGGAAACAGGACTTTTGAGATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 720
DB 100660 AGCCCTCAAACTTTAGAGCGTACATATAGCTTGGAAACAGGACTTTTGAGATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 100719
QY 721 TAAGAAATATCTTTGTAACTCTGAGAACTATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 780
DB 100720 TAAGAAATATCTTTGTAACTCTGAGAACTATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 100779
QY 781 TCAATGAACGTCATCGGCTACAGAAACAGGCTTAAGTTTACAGGACTTTTACAGGCTTTAC 840
DB 100780 TCAATGAACGTCATCGGCTACAGAAACAGGCTTAAGTTTACAGGACTTTTACAGGCTTTAC 100839
QY 841 ATATTTTGGACCAAACTTTGAAGTTGAACCCAGAAACAGACTTTAAACAAATATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 900
DB 100840 ATATTTTGGACCAAACTTTGAAGTTGAACCCAGAAACAGACTTTAAACAAATATATATATATATATATATATATATATATATATATATATATATAGGTTTCT 100899
QY 901 CAATGTATATTTTGTGTGTTTAAAGGCTTGCCTGTGATGTCACAGTTATCCCAATGGAC 960
DB 100900 CAATGTATATTTTGTGTGTTTAAAGGCTTGCCTGTGATGTCACAGTTATCCCAATGGAC 100959
QY 961 ACTAAGTTAGAGCAACAAACCTGATTTCTGGTCTTTCT 1001
DB 100960 ACTAAGTTAGAGCAACAAACCTGATTTCTGGTCTTTCT 101000
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RESULT 2

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AAEF4801
ID AAEF4801 standard; cDNA, 3263 BP.
XX
AC AAEF4801
XX
DB 09-JUL-2001 (first entry)
XX
DE Nucleotide sequence of the human SPG4 polypeptide.
XX
KW Human; SPG4 gene; spastin; PSP-AD; gene therapy;
KW autosomal dominant familial spastic paraplegia; ss.
XX
OS Homo sapiens.
XX
FH Key
CDS 126..1976
FT /*tag= a
FT /product= "spastin"
XX
XX FR2798.38-Al.
XX
PD 09-MAR-2001.
XX
PF 03-SEP-1999; 99FR-0011097.
```

RESULT 3

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ABQ54772
ID ABQ54772 standard; cDNA, 4152 BP.
XX
AC ABQ54772;
XX
DB 22-AUG-2002 (first entry)
XX
DE Human ovarian antigen HNTTB23 cDNA, SEQ ID NO:652.
XX
KW Human; ovarian antigen; ovary; ovarian; breast; cancer; tumour;
KW ovarian cancer; breast cancer; tumour; reproductive system disorder;
KW infertility; pregnancy disorder; anovulation; polycystic ovary syndrome;
XX
```

FR 03-SEP-1999; 99FR-0011097.

XX (CRPS) ; CNRS CENT NAT RECH SCI.

XX Weissentbach J, Hazan J;

XX WP1: 2001-283966/3C.

XX P-PSPB; AAP69137.

XX New human nucleic acid from the SPG4 gene, useful e.g. for diagnosis of autosomal dominant familial spastic paraplegia and in drug screening

XX Claim 2; Page 156-162; 145pp., French.

XX The present sequence encodes a human spastin polypeptide. Spastin is encoded by the SPG4 gene. Mutations in the SPG4 gene are responsible for autosomal dominant familial spastic paraplegia. SPG4 polynucleotides, and their fragments, are used to screen DNA banks for sequences that encode spastin (particularly sequences in other mammals, specifically mice); to identify SPG4 mutations, or other genetic anomalies, particularly for diagnosis of autosomal dominant familial spastic paraplegia (PSP-AD); to identify promoters and other regulatory elements of the SPG4 gene; for detection and amplification; for recombinant production of spastin; and for diagnostic genotyping of PSP-AD.

XX Sequence 3263 BP; 956 A; 664 C; 703 G; 940 T; 0 other;

XX Query Match 40.3%; Score 403; DB 22; Length 3263;

XX Best Local Similarity 100.0%; Pred. No. 8e-64;

XX Matches 403; Conservative 0; Mismatches 0; Gaps 0;

QY 599 AGATGAGAAATATCGATTTATCTGACTTCACTGAATCTGTGAAATAGGTTTCT 658

DB 1952 AGATGAGAAATATCTGATTTATCTGACTTCACTGAATCTGTGAAATAGGTTTCT 1911

QY 659 TCAGCCCTCAAACTTTAGAGCGTACATATAGCTTGGAAACAGGACTTTGGAGATACCACTG 718

DB 1912 TCAGCCCTCAAACTTTAGAGCGTACATATAGCTTGGAAACAGGACTTTGGAGATACCACTG 1971

QY 719 TTTAAGAAATATCTTTGTAACTCTGAGAACTAGGTTTCT 778

DB 1972 TTTAAGAAATATCTTTGTAACTCTGAGAACTAGGTTTCT 2031

QY 779 CTTCAATGAACGTCATCGGCTACAGAAACAGGCTTAAGTTTACAGGACTTTTACAGGCTTTAC 2032

QY 819 ACATATTTTGTGACCAAACTTTGAAGTTGAACCCAGAAACAGACTTTAAACAAATAGGTTTCT 898

DB 2092 ACATATTTTGTGACCAAACTTTGAAGTTGAACCCAGAAACAGACTTTAAACAAATAGGTTTCT 2151

QY 899 TCGAAATGTAAATTTTGTGTTTAAAGGCTTGCCTGTGATGTCACAGTTATCCCAATGG 958

DB 2152 TCGAAATGTAAATTTTGTGTTTAAAGGCTTGCCTGTGATGTCACAGTTATCCCAATGG 2211

QY 919 ACATTAAGTTAGAGCAACAAACCTGATTTCTGGTCTTTCT 1001

DB 2212 ACATTAAGTTAGAGCAACAAACCTGATTTCTGGTCTTTCT 2254

KW PCOS; ovarian cyst; dysmenorrhea; endocrine disorder; infection;
KW inflammatory condition; immune disorder; blood disorder;
KW cardiovascular disorder; respiratory disorder; neurological disorder;
KW gastrointestinal disorder; urinary system disorder; drug screening;
KW gene therapy; chromosome mapping; forensic analysis;
KW antibody preparation; cytostatic; immunomodulatory; neuroprotective;
KW anti-inflammatory; gynaecological; reproductive; gene; ss.
XX
OS Homo sapiens.
XX
XX NC2002200677-A1.
XX
XX 03-JAN-2002.
XX
XX 07-JUN-2001; 2001WC-US18569.
XX
XX 07-JUN-2000; 2000US-209467P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Birse CE, Rosen CA;
XX WPI, 2002-147878/19.
XX P-PSDE; ABP41695.
XX
XX Isolated nucleic acid molecules encoding novel ovarian polypeptides,
XX useful in the prevention, treatment and diagnosis of cancer (e.g.
XX ovarian cancer), immune disorders, cardiovascular disorders and
XX neurological diseases.
XX
XX Claim 1; SEQ ID No 652; 2922bp; English.
XX
XX The invention relates to 2175 novel human ovarian antigens (ABP41654-
XX ABP43228) and to cDNAs encoding them (ABP54331-ABP56335), and also
XX encompasses polypeptides 90% identical, and polynucleotides 95% identical,
XX to the sequences of the invention. The invention additionally relates to
XX recombinant vectors and host cells comprising human ovarian antigen
XX polynucleotides, antibodies against human ovarian antigens, and the use
XX of ovarian antigen polynucleotides and polypeptides in diagnosing,
XX treating, progressing or preventing various ovary and/or breast-related
XX disorders. Such conditions include ovarian cancer and breast cancer, and
XX metastatic tumours of ovarian or breast origin, reproductive system
XX disorders (e.g., infertility, disorders of pregnancy, amenorrhea,
XX polycystic ovary syndrome, ovarian cysts, and dysmenorrhoea), endocrine
XX disorders, infections (e.g., chlamydia, HIV, toxoplasmosis, and toxic
XX shock syndrome), inflammatory conditions (e.g., mastitis, oophoritis and
XX vaginitis), immune disorders (e.g., congenital and acquired
XX immunodeficiencies, autoimmune oophoritis, systemic lupus erythematosus),
XX blood-related disorders (e.g., anaemia), cardiovascular disorders,
XX respiratory disorders, neurological disorders, gastrointestinal disorders
XX and urinary system disorders. Ovarian antigen polypeptides and
XX polynucleotides may also be used in screening for compounds which
XX modulate ovarian antigen expression or activity. The polynucleotides may
XX further be used for gene therapy, chromosome mapping, in the
XX identification of individuals and in forensic analysis, and the
XX polypeptides may be used as food additives or to prepare antibodies
XX useful in disease diagnosis, drug targeting and phenotyping. The present
XX sequence represents cDNA encoding a human ovarian antigen of the
XX invention.
XX Note: The sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pat_sequences.
XX
XX Sequence 4152 BP; 1331 A; 663 C; 777 G; 1376 T; 5 other;
XX
XX Query Match 40.3%; Score 403; DS 24; Length 4152;
XX Best Local Similarity 100.0%; Pred. No. 5,1e-64;
XX Matches 403; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 599 AGATGAGAAATATTCGATTATCTGACTTCACCTGATCCCTGAAAGAAATATAACCCAGCG 658
XX
XX 853 AGATGAGAAATATTCGATTATCTGACTTCACCTGATCCCTGAAAGAAATATAACCCAGCG 912

QY 659 TCAGCCCTCAACTTTAGAACGGTACATACGTTGGAACAGGACTTTGGAGATACCACTG 718
Db 913 TCAGCCCTCAAACTTTAGAACGGTACATACGTTGGAACAGGACTTTGGAGATACCACTG 972
QY 719 TTTAAGGAAATACCTTTTGTAAACCTGCAGAACATTTTACTTTAAAGAGAGSAAACACAGAT 778
Db 973 TTTAAGGAAATACCTTTTGTAAACCTGCAGAACATTTTACTTTAAAGAGAGSAAACACAGAT 1032
QY 779 CTTCAATGAAGCTGATCGGCTACAGAACAGCCCTAAGTTTACAGGACTTTTATAGAGCTTT 838
Db 1033 CTTCAATGAAGCTGATCGGCTACAGAACAGCCCTAAGTTTACAGGACTTTTATAGAGCTTT 1092
QY 839 ACATATTTTGTGACCAAACTTGAAGATGAACAGAACAGACTTTAAACAAAATATACAA 898
Db 1093 ACATATTTTGTGACCAAACTTGAAGATGAACAGAACAGACTTTAAACAAAATATACAA 1152
QY 899 TCCAAATGTAATTTTGTGTTTAAAGCCCTTGGCTTGATGGTCACAGTTATCCCAATGG 958
Db 1153 TCCAAATGTAATTTTGTGTTTAAAGCCCTTGGCTTGATGGTCACAGTTATCCCAATGG 1212
QY 959 ACATTAAGTTAGAGCACACAAACCTGATCTGCTCTCTT 1001
Db 1213 ACATTAAGTTAGAGCACACAAACCTGATCTGCTCTCTT 1255
RESULT 4
AAS17746
ID AAS17746 standard; DNA: 3143 BP.
AC AAS17746;
XX
CT 26-FEB-2002 (first entry)
XX
DE Human genomic clone for P2Y-like G protein-coupled receptor.
XX
KW Human; ds; P2Y-like G protein-coupled receptor; GPCR; GPCR; GPCR;
KW chronic obstructive pulmonary disease; nervous system disease;
KW Parkinson's disease; multiple sclerosis; dementia; stroke;
KW Alzheimer's disease; benign prostatic hyperplasia; urinary incontinence;
KW bacterial infection; fungal infection; protozoan infection;
KW viral infection; pain; cancer; anorexia; bulimia; asthma;
KW acute heart failure; hypertension; hypertension; osteoporosis; diabetes;
KW angina pectoris; myocardial infarction; ulcer; inflammation; allergy;
KW psychotic disorder; neurological disorder; anxiety; schizophrenia;
KW manic depression; delirium; severe mental retardation; dyskinesia.
XX
CS Homo sapiens.
XX
XX Key Location/Qualifiers
XX CDS 520..2498
XX /*tag= d
XX /product= "P2Y-like GPCR"
XX
XX NC200185754-A2.
XX
XX 15-NOV-2001.
XX
XX 09-MAY-2001; 2001WC-EP05244.
XX
XX 11-MAY-2000; 2000US-203582P.
XX 21-FEB-2001; 2001US-269857P.
XX
XX (PAB) : BAYER AG.
XX
XX Ramakrishnan S;
XX
XX WPI; 2002-075242/10.
XX P-PSDE; AAU11251.
XX
XX New polynucleotides for producing P2Y-like G protein-coupled receptors
XX (GPCR) that are used for screening inhibitors or regulators of human
XX P2Y-like GPCR, especially useful for treating pain, cancer or
XX neurological disorders.

XX Disclosure: Fig 1: 114pp; English.
 XX
 XX
 CC The invention relates to an isolated polynucleotide encoding a p2Y-like
 CC G protein-coupled receptor (GPCR), polypeptide, its fragment,
 CC derivative or allele, a host cell containing an expression vector
 CC comprising the polynucleotide and screening for agents that regulate the
 CC GPCR activity. The polynucleotide is useful for producing p2Y-like GPCR
 CC polypeptide, which may be employed for screening agents that inhibit or
 CC regulate human p2Y-like GPCR. The reagent or inhibitor of the human
 CC p2Y-like GPCR is useful for treating or ameliorating p2Y-like GPCR
 CC disorders, particularly COPD, chronic obstructive pulmonary disease,
 CC peripheral or central nervous system disease (e.g., Parkinson's
 CC disease, multiple sclerosis, dementia, stroke, Alzheimer's disease and
 CC many other diseases and disorders listed in the specification), benign
 CC prostatic hyperplasia or urinary incontinence, A pharmaceutical
 CC composition containing the modulators and/or regulators of p2Y-like
 CC GPCR is useful for modulating the activity of a p2Y-like GPCR.
 CC In particular, these are useful for treating, preventing or ameliorating
 CC infections (e.g., bacteria, fungi, protozoan or viral infections), pain,
 CC cancer, anorexia, bulimia, asthma, acute heart failure, hypotension,
 CC hypertension, osteoporosis, diabetes, angina pectoris, myocardia,
 CC infarction, ulcers, inflammation, allergies, psychotic or neurologial
 CC disorders (e.g., anxiety, schizophrenia, manic depression, delirium,
 CC severe mental retardation or dyskinesias). The present sequence is
 CC a genomic clone encoding the p2Y-like GPCR of the invention.
 XX
 XX Sequence 3-43 BP; 818 A; 77 C; 726 G; 828 T; 0 other;
 SQ
 Query Match 19.1%; Score 191.4; DB 24; Length 3143;
 Best Local Similarity 88.8%; Pred. No. 9.9e+26;
 Matches 207; Conservative 0; Mismatches 26; Indels 0; Gaps 0;
 QY 13 CATACTTAGGCTGGGCAAGTGGCTCACCTCTGTATCCAGCACTTTGGGAGCCCAAG 72
 Db 2723 CATATCTAGGCTGGGCACATGGCTCATGCTGTATCCAGCACTTTGGGAGCCCAAG 2782
 QY 73 GTGGCGGATCACCTGAGGTTAGGAGTTGAGGACCAACCTGGCCATCGGCAAAACCT 132
 Db 2763 GGCGGGGATCACCTGAGGTTGAGGAGTTGAGGACCAACCTGGCCAACTGGTGAACCCG 2842
 QY 133 GGCTCTACTAAANATACAAAANTTAGCTGGGCATGGAGGCACTGCCTGTATCCCACT 192
 Db 2843 ATCTCTACTAAANATACAAAANTTAGCTGGGCGGTGGTGGCGGTGCTATATCCCACT 2922
 QY 193 ACTCGGAGGTTGACAGAGAGAAATTCCTTGAACCTTGGAGGCGGAGGTGCA 243
 Db 2903 ACTCGGAGGCTTGAGAAAGAGAAATTCCTTGAACCGCGGAGCGGAGGTGCA 2955
 RESULT 5
 AA104191
 ID AAL04181 standard; DNA; 30573 BP.
 XX
 XX AAL04181;
 XX
 XX 21-NOV-2001 (first entry)
 XX
 XX Human reproductive system related antigen DNA SEQ ID NO: 4969.
 DE
 XX Human reproductive system related antigen; reproductive system disorder;
 XX cancer; gene therapy; ds.
 XX
 XX Homo sapiens.
 OS
 XX
 XX WC020153120-A2.
 PN
 XX
 XX 02-AUG-2001.
 XX
 XX 17-JAN-2001; 2001WO-US01339.
 XX
 XX 31-JAN-2000; 2000US-0179065.
 PR
 XX 04-FEB-2000; 2000US-0180626.


```
Db 12886 TAAATAACAAATTTAGCTGGGCAATGGTGGCAGCTGCTGTAAATCCCGACTCTCTGGGA 12945
QY 201 GGCTGAGACAGAGAAATTCCTTGAACCTTGGAGGGGAGGTTGCA 245
Db 12946 GGCTGAGGAGGAGAAATTCCTTGAACCTTGGAGGGGAGGTTGCA 12990

RESULT 7
ABV23205
ID ABV23205 standard; cDNA: 7677 BP.
XX AC ABV23205;
XX DT 16-SEP-2002 (first entry)
DE Human prostate expression marker cDNA 23196.
XX KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
XX KW pharmacogenomic marker; gene; ss.
XX OS Homo sapiens.
XX PN WC200162860-A2.
XX PD 23-AUG-2001.
XX PF 23-FEB-2001; 2001WC-USC5171.
XX PS 17-FEB-2000; 2000US-183319P.
XX PR 16-MAR-2000; 2000US-189862P.
XX PR 25-MAY-2000; 2000US-227454P.
XX PR 09-JUN-2000; 2000US-211314P.
XX PR 18-JUL-2000; 2000US-219037P.
XX PR 13-DEC-2000; 2000US-255281P.
XX PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
XX X Schlegel R, Endege WO, Morahan JE;
XX WPI; 2001-662795/76.
XX PT Novel isolated nucleic acid molecule associated with cancerous state of
XX PT prostate cells and correlating with presence of prostate cancer, useful
XX PT for detecting presence of prostate cancer, stage of prostate cancer.
XX PS Claim 1; Page 4173-4175; 11750pp; English.
XX CC The invention relates to an isolated nucleic acid molecule (1) comprising
XX CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV52213) of the
XX CC specification or its complement. (1) is useful for:
XX CC (a) assessing whether a patient is afflicted with prostate cancer;
XX CC (b) monitoring the progression of prostate cancer in a patient;
XX CC (c) assessing the efficacy of a test compound to inhibit prostate
XX CC cancer in a patient;
XX CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer
XX CC in a patient;
XX CC (e) selecting a composition for inhibiting prostate cancer in a patient;
XX CC (f) assessing the prostate cell carcinogenic potential of a compound;
XX CC (g) determining whether prostate cancer has metastasized in a patient;
XX CC (h) assessing the aggressiveness or incidence of prostate cancer in a
XX CC patient;
XX CC (1) is also useful as a pharmacodynamic or pharmacogenomic marker.
XX X Sequence 7677 BP; 2389 A; 1642 C; 1794 G; 1846 T; 6 other;
S2

Query Match 19.1%; Score 190.8; DB 23; Length 7677;
Best Local Similarity 60.7%; Pred. No. 1.3e-25;
Matches 368; Conservative 0; Mismatches 227; Indels 11; Gaps 3;

QY 1 AATATAAATGCACTACTTAGCTGGGCAAGTGGCTCAGCTGTGTAATCCCGACTT 60
Db 6036 AATTTTAAAGAAATTAATCTCGGCTGGGCGGCTGGCTCAGCTGTGTAATCCCGACTT 6095
```

```
QY 61 TGGAGGCGCAAGSTGGGCGGATCACTTGAGGTTAGGAGTTGAGGCAAACTTGGCCATCA 120
Db 6896 TGGGAAGCGCAAGGTGGGCGGATCACTTTCAGGTTAGGAGTTGAGGACAGGCTTGGCAACA 6955
QY 121 TGGCGAAAACCCCTGCTCTACTTAAATAATACAAAAATAGCTGGGCAATGGAGGCATGTGCTT 180
Db 6956 TGGCGAAAACCCCTGCTCTACTTAAATAATACAAAAATAGCCGGGGCTGGTGGCACTTGCCT 7015
QY 181 GTAATCCAGCTACTCGGAGGCTGAGACAGAGAAATTCGTTGAACCTTGGAGGGGAGG 240
Db 7016 GTAATCCAGCTACTCGGAGGCTGAGGACAGAGAAATTCGATTAACCTGGGAGCGGAGA 7075
QY 241 TTGC---ATACTGAGTGGTGAATGTGATTCCTTTTCTCTTTGCTGATTTTGG 296
Db 7076 TTGCAATGAGTGGAGCCACTGCACTCGGCTGGGTGACAGAGAGAGACTGCTCTGG 7135
QY 297 AACTTTTCATAAATGATTGTTGTTTATATTTTATATTTGGAATAATATATGCTTTCAAT 356
Db 7136 AAAAAAAGAAATCTCACTCACTATCTAGAGAGGATTGTCAGAAATATTCACGATTCAGT 7195
QY 357 GTTAATACCTATGAACCTAAACAGAGTAAATAATATAGTATAGTATAGCACTTATTAGGT 416
Db 7196 CTTGAAACTTTGATTATGCAAAAGAGTATATATATAATTTTCAATATGATTCAGTTT 7255
QY 417 TTCTTGTGACAGATCAACATAGAAATATATTAATGGCTGACATATTTTCTAAGA 476
Db 7256 TTAAGGCTTTCAGCT---CTATAAGTGTCTCAGATGCCACTAGATAATTTTAAAGC 7312
QY 477 ATACATACAGCTATATTTTATACATTA---AGAAAGAGGAGCATATTACTTTAAT 532
Db 7313 ATCATATTAGAAATACTTTAAGAGACTTATATAGAAATAGAAAGATGTTGAAATTTAC 7372
QY 533 GCATCATTTGGTTACCAACCATATACCTGTTGATCATTTGATTTGATGTCATGCTTTTAA 592
Db 7373 AGAGGATTTGGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTT 7432
QY 593 AATCT 595
Db 7433 ACATAT 7438

RESULT 8
ABV29042
ID ABV29042 standard; cDNA: 7677 BP.
XX AC ABV29042;
XX CX 16-SEP-2002 (first entry)
XX DE Human prostate expression marker cDNA 29033.
XX XX Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
XX KW pharmacogenomic marker; gene; ss.
XX OS Homo sapiens.
XX XX WC200162860-A2.
XX PD 23-AUG-2001.
XX PF 23-FEB-2001; 2001WC-USC5171.
XX PR 17-FEB-2000; 2000US-183319P.
XX PR 16-MAR-2000; 2000US-189862P.
XX PR 25-MAY-2000; 2000US-227454P.
XX PR 09-JUN-2000; 2000US-211314P.
XX PR 18-JUL-2000; 2000US-219037P.
XX PR 13-DEC-2000; 2000US-255281P.
XX PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
XX X Schlegel R, Endege WO, Morahan JE;
XX P;
```

XX W21: 2001-662795/76.
XX Novel isolated nucleic acid molecule associated with cancerous state of
XX prostate cells and correlating with presence of prostate cancer, useful
XX for detecting presence of prostate cancer, stage of prostate cancer
XX Claim 1: Page 6152-6154; 1:750pp; Engl.sh.
XX The invention relates to an isolated nucleic acid molecule (I) comprising
XX a nucleotide sequence given in Tables 1-9 (ABV30010-ABV62233) of the
XX specification or its complement (II), is useful for:
XX (a) assessing whether a patient is afflicted with prostate cancer;
XX (b) monitoring the progression of prostate cancer in a patient;
XX (c) assessing the efficacy of a test compound to inhibit prostate
XX cancer in a patient;
XX (d) assessing the efficacy of a therapy for inhibiting prostate cancer
XX in a patient;
XX (e) selecting a composition for inhibiting prostate cancer in a patient;
XX (f) assessing the prostate cell carcinogenic potential of a compound;
XX (g) determining whether prostate cancer has metastasized in a patient;
XX (h) assessing the aggressiveness or indolence of prostate cancer in a
XX patient;
XX (i) is also useful as a pharmacodynamic or pharmacogenomic marker.
XX Sequence 7677 BP; 2389 A; 1642 C; 1794 G; 1846 T; 6 other;
XX
XX Query Match: 19.1%; Score 190.8; DB 23; Length 7677;
XX Best Local Similarity 60.7%; Pred. No. 1.3e-25;
XX Matches 368; Conservative 0; Mismatches 227; Indels 11; Gaps 3;
XX
XX 1 AATAAATAATGCACTTATAGGTGGCAAGTGGCTCAGCTGTGTAAATCCAGACACT 60
XX 5836 AATTATTAAGNAATTAATCTCGGTGGCGCGGTGGCTCAGCTGTGTAAATCCAGACACT 6895
XX
XX 61 TGGGAGGCGCAAGTGGCGGATCACTCAGGTAGGAGTTCAGGACCAACTGGCCATCA 120
XX 6896 TGGGAGGCGCAAGTGGCGGATCACTCAGGTAGGAGTTCAGGACCAACTGGCCATCA 6955
XX
XX 121 TGGGAGGCGCGTCTTACTAATAATACAAATATAGCTGGGATGGAGGATGGCT 180
XX 6956 TGGGAGGCGCGTCTTACTAATAATACAAATATAGCTGGGATGGAGGATGGCT 7015
XX
XX 181 GTAAATCCAGCTACTCGGAGGCTGAGACAGAAATTTGTTGAACCTTGGAGGGGAGG 240
XX 7016 GTAAATCCAGCTACTCGGAGGCTGAGACAGAAATTTGTTGAACCTTGGAGGGGAGG 2475
XX
XX 241 TTGC-----ATATCTAGTGGTGAATTTGATTTCTTTTCTCTCTTCTGCTGATTTTG 236
XX 7076 TTGCAATGAGATCGAACCACTGGCACTCGAGCTGGTGGAGAGAGAGACCTGGCTTG 7135
XX
XX 297 AACTTTTCTATAAATGATGTTGTTTGTGTTTATATATGGAANAATATATGCTTCAAT 336
XX 7136 AAAAAAAGAAATGTCACCTACTATCTAGAGAGAGATTCAGAAATATTCAGGATTCAGGT 7195
XX
XX 357 GTATATACCTATGAACCTAATACACAGTAATAAATAATATAGTATAGTATTTATAGGT 416
XX 7196 CTTGAAACCTTGAATATGCAAAAGAGGTATATATATAATATATATATATATATATAT 7255
XX
XX 417 TCTCTGTGTAGCAGATCAACATAGAAATATATATTAATGGCTGACATATTTCTAAGA 476
XX 7256 TTAGGCTTTGAGCT---TCTATAGTGTCTTCTAGATGGCACTAGATATTTTAAAGC 7312
XX
XX 477 ATACATACAGGTATATTTTATAACATTA----AGAAACAGGAGCTATATTTACTTAAT 532
XX 7313 ATCAATTAAGAAATACTTTAAGAGACCTATATAAGAAATACAGAGATTTGGAATTTAC 7372
XX
XX 533 CAGTATCTCGTATACCACTATACCTGTTGATCAATTTGTATCTGATGCTGCTTTTAA 592
XX 7373 AGAGAGATTGGTTCATTAAGAGCCAGATTCGTAAAGTTTTCATTTGAAATTTTAGCTAA 7432
XX
XX 593 AAATCT 538
XX | | |

Db 7433 ACATAT 7438
RESULT 9
AAC76994/C
ID AAC76994 standard, CONA; 1074 BP.
XX AAC76994;
XX AAC76994;
XX 09-FEB-2001 (first entry)
DE Hu-an ORFX ORF549 polynucleotide sequence SEQ ID NO:5097.
XX
XX Human; open reading frame; ORFX; detection; cytostatic; hepatotropic;
XX valvular; antiproliferative; antiparkinsonian; nootropic; neuroprotective;
XX anticonvulsant; osteopathic; antiarthritic; immunosuppressant; cardiac;
XX immunostimulant; thrombolytic; coagulant; vasotropic; antidiabetic;
XX hypotensive; dermatological; immunosuppressive; antiinflammatory;
XX antiviral; antibacterial; antifungal; antipneumatic; antithyroid;
XX antianemic; gene therapy; cancer; proliferative disorder; hypertension;
XX neurodegenerative disorder; osteoarthritis; graft vs host disease;
XX cardiovascular disease; diabetes mellitus; hypothyroidism; SCID; AIDS;
XX cholesterol ester storage; systemic lupus erythematosus; infection;
XX severe combined immunodeficiency; malaria; autoimmune disorders; asthma;
XX allergy; aplastic anaemia; nocturnal haemoglobinuria; burn; wound;
XX bone damage; cartilage damage; antiinflammatory disease; coagulation;
XX thrombosis; contraceptive; ss.
XX
XX Homo sapiens.
XX WC200358473-A2.
XX
XX 05-OCT-2000.
XX
XX 31-MAR-2000; 2000WO-US08621.
XX
XX 31-MAR-1999; 99US-0127607.
XX 02-APR-1999; 99US-0127636.
XX 05-APR-1999; 99US-0127728.
XX 30-MAR-2000; 2000US-0340763.
XX
XX (CURA-); SURAGEN CORP.
XX
XX Shinketsu RA, Leach M;
XX WPI: 2000-602662/57.
XX P-PSDB: AAB42765.
XX
XX Novel nucleic acids and peptides derived from open reading frame X,
XX useful for treating e.g. cancers, proliferative disorders,
XX neurodegenerative disorders and cardiovascular disease -
XX
XX Claim 5: Page 4285-4287; 5507pp; English.
XX
XX AAC769446 to AAC76960 encode the proteins given in AAB40237 to AAB43397,
XX which represent the human ORFX open reading frames 1 to 3161. The ORFX
XX sequences have activities such as: cytostatic; hepatotropic; valvular;
XX antiproliferative; antiparkinsonian; nootropic; neuroprotective;
XX osteopathic; anticonvulsant; antiarthritic; immunosuppressant;
XX immunostimulant; cardiac; thrombolytic; coagulant; vasotropic;
XX antidiabetic; hypotensive; dermatological; immunosuppressive;
XX antiinflammatory; antibacterial; antiviral; antifungal; antipneumatic;
XX antithyroid; and antianemic. The sequences can be used for determining
XX the presence of or predisposition to, or preventing or treating
XX pathological conditions associated with an ORFX-associated disorder. The
XX nucleic acids can be used to express ORFX proteins in gene therapy.
XX vectors. The proteins and nucleic acids may be used to treat cancers,
XX proliferative disorders, neurodegenerative disorders, osteoarthritis,
XX graft vs host disease, cardiovascular disease, diabetes mellitus,
XX hypertension, hypothyroidism, cholesterol ester storage, systemic lupus
XX erythematosus, severe combined immunodeficiency (SCID), AIDS, viral,
XX bacterial or fungal infection, malaria, autoimmune disorders, asthma,
XX allergies, aplastic anaemia, burns, wounds, bone and cartilage damage,
XX

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CC nocturnal haemoglobinuria, antiinflammatory disease; to enhance
CC coagulation; to inhibit thrombosis; and as a contraceptive.
XX
SQ Sequence 3074 BP; 654 A; 851 C; 847 G; 720 T; 2 other;

Query Match 19.0%; Score 190.4; DB 21; Length 3074;
Best Local Similarity 87.1%; Pred. No. 1.5e-25;
Matches 209; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

QY 6 AAAAATGATCTATTAGCTGGGCAAGTGGCTCACGTTCTTAATCCAGCACATTGGGA 65
DB 1593 AAAAGTGCTTTTAGAGGCTGGCATGGTGCTCAGGCTTGAATCCAGCACATTGGGA 1594
QY 66 GCGCAGGTGGGCGGATCACTTGAGGTTAGAGTTAGGACCAACCTGGCCATCATGGCG 125
DB 1533 GGCCAAGGTGGGTGGATCACTTGAGGTTAGAGTTAGGACCAACCTGGCCATCATGGTG 1474
QY 126 AAACCTCTCTCTACTAAATAAGAAATAGCTGGCATGGGATGGCTGTGAAT 185
DB 1473 AAACCTCTCTCTACTAAATAAGAAATAGGACGAGCAATGGTGTGAAT 1474
QY 186 CCCAGCTACTCGGAGGCTGAGACAGAAAGAAATTCCTGAACCTTGGAGGGAGGTTTSCA 245
DB 1413 CCCAGCTACTGGGAAGCTGAGGACGAGAAATTCCTGAACCTTGGAGGGAGGTTTSCA 1354

RESULT 10
AAAT67164
ID AA*67164 standard; cDNA; 10380 BP.
XX
AC AA*67164;
CX 20-AUG-1997 (first entry)
DE Human alpha-N-acetylglucosaminidase gene.
KW Alpha-N-acetylglucosaminidase; mucopolysaccharidosis type IIIB;
KW gene therapy; enzyme replacement therapy; diagnosis; ss.
XX Homo sapiens.
FH Key Location/Qualifiers
FT 5'UTR 1..989
FT exon /*tag= a
FT 990..1372
FT /*tag= b
FT intron /*tag= c
FT 1373..2214
FT exon /*tag= d
FT 2115..2262
FT /*tag= e
FT intron /*tag= f
FT 2263..3055
FT intron /*tag= g
FT 3056..3202
FT /*tag= h
FT intron /*tag= i
FT 3203..3386
FT exon /*tag= j
FT 3387..3472
FT intron /*tag= k
FT 3473..5666
FT exon /*tag= l
FT 5667..5923
FT intron /*tag= m
FT 5924..7744
FT exon /*tag= n
FT 7745..8955
FT intron /*tag= o
FT 8956..10380
FT 3'UTR

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FT XX WC9719177-A1. /*tag= n
XX XX 29-MAY-1997.
XX XX 22-NOV-1996; 96WO-AU00747.
XX XX 23-NOV-1995; 95AU-0006748.
XX XX (WOMEN'S & CHILDREN'S HOSPITAL.
XX XX Anson DS, Blanch L, Hopwood JJ, Scott H, Weber B;
XX MPI; 1997-298:14/27.
XX DR P-PSDB; AAW18017.
XX PT Nucleic acid encoding mammalian alpha-N-acetylglucosaminidase
XX used for the diagnosis and treatment of mucopolysaccharidosis type
XX IIIB, also used in gene therapy
XX
XX Claim 3; Page 54-61; 79pp; English.
XX
XX A genomic DNA molecule (AA*67164) includes 6 exons that code for
XX human alpha-N-acetylglucosaminidase (AAW18017), an enzyme that can
XX hydrolyse the terminal alpha-N-acetylglucosamine residues at the
XX non-reducing terminus of fragments of heparan sulphate and heparin.
XX It was isolated by hybridisation of a human chromosome 17 library.
XX A cDNA clone (AA*67163) coding for the enzyme has also been isolated.
XX The isolated gene or cDNA, and primers/probes based on them or
XX their complementary strands, can be used to investigate, diagnose
XX and treat alpha-N-acetylglucosaminidase deficiency, for example in
XX patients suffering from mucopolysaccharidosis type IIIB.
XX Administration is by oral, i.v., i.p., enzyme replacement therapy,
XX gene therapy or other routes.
XX
XX Sequence 10380 BP; 2210 A; 2953 C; 2851 G; 2366 T; 0 other;
XX
Query Match 19.0%; Score 189.8; DP 15; Length 10380;
Best Local Similarity 86.7%; Pred. No. 2e-25;
Matches 209; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

QY 5 TAAATATGATCTATTAGCTGGGCAAGTGGCTCACGTTCTTAATCCAGCACATTGGG 64
DB 3540 TAAATATGATCTATTAGCTGGGCGGATGGCTCACGTTCTTAATCCAGCACATTGGG 3599
QY 65 AGCCCAAGTGGGCGGATCACTTGAGGTTAGGAGTTAGGAGCAACCTGGCCATCATGGC 124
DB 3600 AGCCCGAGTTGGGCGGATCACTTGAGGTTAGGAGTTAGGAGCAACCTGGCCATCATGGT 3659
QY 125 GAAACCTGTCTCTACTTAAATAAGAAATAGCTGGCATGGGAGGATGTGCTGTGA 184
DB 3660 GAAACCTGTCTCTACTTAAATAAGAAATAGCTGGCATGGGAGGATGTGCTGTGA 3719
QY 185 TCCAGCTACTCGGAGGCTGAGACAGAAAGAAATTCCTGAACCTTGGAGGGAGGTTGC 244
DB 3720 TCCAGCTACTTAGGAGGCTGAGACAGGAGAAATTCCTGAACCTTGGAGGGAGGTTGC 3779
QY 245 A 245
DB 3780 A 3780

RESULT 1:
AAS32727
ID AAS32727 standard; DNA; 17792 BP.
XX
AC AAS32727;
XX XX 17-DEC-2001 (first entry)
XX Human genomic DNA for novel endocrine antigen, SEQ ID No 681.
XX

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PR 01-DEC-2000; 2000US-0251160.
PR 05-DEC-2000; 2000US-0251032.
PR 05-DEC-2000; 2000US-0251988.
PR 05-DEC-2000; 2000US-0256719.
PR 06-DEC-2000; 2000US-0251479.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.
XX

PA (HUMA-) HUMAN GENOME SC: INC.

XX Rosen CA, Barash SC, Ruben SM;

XX WPI; 2001-541565/60.

XX Nucleic acids encoding 3224 human nervous system antigen polypeptides,
PT useful for preventing, diagnosing and/or treating nervous system
PT cancers and metastases.

XX Disclosure, SEQ ID NO 8425; 1701pp + Sequence Listing; English.

XX The invention relates to novel genes (ABA11004-ABA21534) and proteins
CC (ABA14678-ABA19301) useful for preventing, treating or ameliorating
CC medical conditions e.g. by protein or gene therapy. The genes are
CC isolated from a range of human tissues disclosed in the specification.
CC The nucleic acids, proteins, antibodies and (ant)agonists are useful
CC in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast
CC and ovarian cancer and other cancers of the adrenal gland, bone, bone
CC marrow, breast, gastrointestinal tract, liver, lung, CR urogenital;
CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune
CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative
CC colitis; (c) cardiovascular disorders such as myocardial ischaemias;
CC (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and
CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal
CC and parasitic infections.

CC Note: The specification data for this patent did not form part of the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

XX Sequence 15616 BP; 4181 A; 5695 C; 5539 G; 4201 T; 0 other;

Query Match: 18.8%; Score 188.6; DB 22; Length 19616;
Best Local Similarity 86.0%; Pred. No. 3,48,23;
Matches 209; Conservative 0; Mismatches 34; Indels 0; Gaps 0;
QY 3 TATAAAATGCACTTTAGCTGCGGCAAAATGGCTCAGCTCTGTATATCCAGCACTTG 62
DB 17028 TGTGTAAATGCTAAATGGGSCCAAGCGCACTGCTCAGCGCTTATATCCAGCACTTG 16969
QY 63 GGAGCGGAGGTGGGCGGATACCTGAGGTAGGATTCAGGACCACTTGGGCGCTCAG 122
DB 16969 GGAGCGGAGGTGGGCGGATACCTGAGGTAGGATTCAGGACCACTTGGGCGCACTG 16909
QY 123 GGGAACCTCTCTCTACTATAAATACAAAATAGTGGGCAATGAGGCGATATGCTCT 182
DB 16938 ACGAAACCTCTCTCTACTATAAATACAAAATAGTGGGCGGCTGGTGGCACTGCT 16843
QY 183 ATCCAGCTATCCGGAGGCTGAGACAGAGCAATTCCTTGAAGCTTGGAGGGGAGCTT 212
DB 16848 ATCCAGCTATCCGGAGGCTGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 16789
QY 243 GCA 245
DB 16788 GCA 16786

RESULT 14
ABA16856/c

IC ABA16856 standard; DNA; 19616 BP.
XX
XX ABA16856;
XX
XX 23-JAN-2002 (first entry)
XX Human nervous system related polynucleotide SEQ ID NO 9187.
XX
XX Human; neurotropic; neuroprotective; cytostatic; dermatological; virucide;
XX immunosuppressive; antiinflammatory; anti-HIV; antibacterial; vulvovag;
XX antiparkinsonian; antiskilling; antianemic; antituberc; cancer;
XX antituberc; hepatotropic; cerebroprotective; antiinflammatory;
XX anti-herpetic; antidiabetic; antitumor; anticonvulsant; antifungal;
XX antiparasitic; cardiac; immune disorder; cardiovascular disorder;
XX neurological disease; infection; nephrotropic; gene therapy; vaccine; ds.
XX
XX HOTA sapiens.
XX
XX WO200159563-A2.
XX
XX 16-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US01334.
XX
XX 31-JAN-2000; 2000US-0179565.
XX 04-FEB-2000; 2000US-0180528.
XX 24-FEB-2000; 2000US-0184664.
XX 02-MAR-2000; 2000US-0186350.
XX 16-MAR-2000; 2000US-0189874.
XX 17-MAR-2000; 2000US-0190076.
XX 18-APR-2000; 2000US-0198123.
XX 19-MAY-2000; 2000US-0205515.
XX 07-JUN-2000; 2000US-0209467.
XX 28-JUN-2000; 2000US-0214886.
XX 30-JUN-2000; 2000US-0225135.
XX 07-JUL-2000; 2000US-0216647.
XX 07-JUL-2000; 2000US-0216880.
XX 11-JUL-2000; 2000US-0217487.
XX 11-JUL-2000; 2000US-0217496.
XX 14-JUL-2000; 2000US-0218290.
XX 26-JUL-2000; 2000US-0220963.
XX 26-JUL-2000; 2000US-0220964.
XX 14-AUG-2000; 2000US-0224518.
XX 14-AUG-2000; 2000US-0224519.
XX 14-AUG-2000; 2000US-0225133.
XX 14-AUG-2000; 2000US-0225214.
XX 14-AUG-2000; 2000US-0225266.
XX 14-AUG-2000; 2000US-0225267.
XX 14-AUG-2000; 2000US-0225268.
XX 14-AUG-2000; 2000US-0225270.
XX 14-AUG-2000; 2000US-0225447.
XX 14-AUG-2000; 2000US-0225757.
XX 14-AUG-2000; 2000US-0225758.
XX 14-AUG-2000; 2000US-0225759.
XX 18-AUG-2000; 2000US-0226279.
XX 22-AUG-2000; 2000US-0226681.
XX 22-AUG-2000; 2000US-0226868.
XX 22-AUG-2000; 2000US-0227182.
XX 23-AUG-2000; 2000US-0227639.
XX 30-AUG-2000; 2000US-0228924.
XX 01-SEP-2000; 2000US-0229287.
XX 01-SEP-2000; 2000US-0229343.
XX 01-SEP-2000; 2000US-0229344.
XX 01-SEP-2000; 2000US-0229345.
XX 05-SEP-2000; 2000US-0229509.
XX 05-SEP-2000; 2000US-0229513.
XX 06-SEP-2000; 2000US-0230437.
XX 06-SEP-2000; 2000US-0230438.
XX 08-SEP-2000; 2000US-0231242.
XX 08-SEP-2000; 2000US-0231243.
XX 08-SEP-2000; 2000US-0231244.
XX 08-SEP-2000; 2000US-0231413.
XX 08-SEP-2000; 2000US-0231414.
XX

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PR 08-SEP-2000; 2000US-023280.
PR 08-SEP-2000; 2000US-023281.
PR 12-SEP-2000; 2000US-0231968.
PR 14-SEP-2000; 2000US-0231969.
PR 14-SEP-2000; 2000US-023197.
PR 14-SEP-2000; 2000US-023198.
PR 14-SEP-2000; 2000US-023199.
PR 14-SEP-2000; 2000US-0232400.
PR 14-SEP-2000; 2000US-0232401.
PR 14-SEP-2000; 2000US-0231063.
PR 14-SEP-2000; 2000US-0231064.
PR 14-SEP-2000; 2000US-0231065.
PR 21-SEP-2000; 2000US-0234223.
PR 21-SEP-2000; 2000US-0234274.
PR 25-SEP-2000; 2000US-0234997.
PR 25-SEP-2000; 2000US-0234998.
PR 26-SEP-2000; 2000US-0234984.
PR 27-SEP-2000; 2000US-0235834.
PR 27-SEP-2000; 2000US-0235836.
PR 29-SEP-2000; 2000US-0236327.
PR 29-SEP-2000; 2000US-0236367.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236370.
PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
PR 02-OCT-2000; 2000US-0237038.
PR 02-OCT-2000; 2000US-0237039.
PR 02-OCT-2000; 2000US-0237042.
PR 13-OCT-2000; 2000US-0239335.
PR 13-OCT-2000; 2000US-0239337.
PR 20-OCT-2000; 2000US-0240560.
PR 20-OCT-2000; 2000US-0241785.
PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 20-OCT-2000; 2000US-0242221.
PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
PR 08-NOV-2000; 2000US-0246478.
PR 08-NOV-2000; 2000US-0246523.
PR 08-NOV-2000; 2000US-0246524.
PR 08-NOV-2000; 2000US-0246525.
PR 08-NOV-2000; 2000US-0246526.
PR 08-NOV-2000; 2000US-0246527.
PR 08-NOV-2000; 2000US-0246611.
PR 08-NOV-2000; 2000US-0246613.
PR 17-NOV-2000; 2000US-0249237.
PR 17-NOV-2000; 2000US-0249238.
PR 17-NOV-2000; 2000US-0249239.
PR 17-NOV-2000; 2000US-0249210.
PR 17-NOV-2000; 2000US-0249211.
PR 17-NOV-2000; 2000US-0249212.
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PR 17-NOV-2000; 2000US-0249214.
PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249299.
PR 08-SEP-2000; 2000US-023280.
PR 08-SEP-2000; 2000US-023281.
PR 12-SEP-2000; 2000US-0231968.
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PR 14-SEP-2000; 2000US-023198.
PR 14-SEP-2000; 2000US-023199.
PR 14-SEP-2000; 2000US-0232400.
PR 14-SEP-2000; 2000US-0232401.
PR 14-SEP-2000; 2000US-0231063.
PR 14-SEP-2000; 2000US-0231064.
PR 14-SEP-2000; 2000US-0231065.
PR 21-SEP-2000; 2000US-0234223.
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PR 27-SEP-2000; 2000US-0235836.
PR 29-SEP-2000; 2000US-0236327.
PR 29-SEP-2000; 2000US-0236367.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236370.
PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
PR 02-OCT-2000; 2000US-0237038.
PR 02-OCT-2000; 2000US-0237039.
PR 02-OCT-2000; 2000US-0237042.
PR 13-OCT-2000; 2000US-0239335.
PR 13-OCT-2000; 2000US-0239337.
PR 20-OCT-2000; 2000US-0240560.
PR 20-OCT-2000; 2000US-0241785.
PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 20-OCT-2000; 2000US-0242221.
PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
PR 08-NOV-2000; 2000US-0246478.
PR 08-NOV-2000; 2000US-0246523.
PR 08-NOV-2000; 2000US-0246524.
PR 08-NOV-2000; 2000US-0246525.
PR 08-NOV-2000; 2000US-0246526.
PR 08-NOV-2000; 2000US-0246527.
PR 08-NOV-2000; 2000US-0246611.
PR 08-NOV-2000; 2000US-0246613.
PR 17-NOV-2000; 2000US-0249237.
PR 17-NOV-2000; 2000US-0249238.
PR 17-NOV-2000; 2000US-0249239.
PR 17-NOV-2000; 2000US-0249210.
PR 17-NOV-2000; 2000US-0249211.
PR 17-NOV-2000; 2000US-0249212.
PR 17-NOV-2000; 2000US-0249213.
PR 17-NOV-2000; 2000US-0249214.
PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249299.

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17-NOV-2000; 2000US-0249300.
 21-DEC-2000; 2000US-0250191.
 21-DEC-2000; 2000US-0251150.
 25-DEC-2000; 2000US-0251030.
 25-DEC-2000; 2000US-0251988.
 25-DEC-2000; 2000US-0256719.
 26-DEC-2000; 2000US-0251479.
 28-DEC-2000; 2000US-0251856.
 28-DEC-2000; 2000US-0251868.
 28-DEC-2000; 2000US-0251869.
 28-DEC-2000; 2000US-0251989.
 28-DEC-2000; 2000US-0251990.
 11-DEC-2000; 2000US-0254097.
 05-JAN-2001; 2001US-0259678.
 (HUMA-) HUMAN GENOME SCI INC.
 Rosen CA, Barash SC, Ruben SM;
 WPI; 2001-541565/60.
 Nucleic acids encoding 3224 human nervous system antigen polypeptides,
 useful for preventing, diagnosing and/or treating nervous system
 cancers and metastases.
 Disclosure; SEQ ID NO 9-97; 1701bp - Sequence listing; English.
 The invention relates to novel genes (ABA1004-ABA21534) and proteins
 (ABP4578-ABP1601) useful for preventing, treating or ameliorating
 medical conditions e.g. by protein or gene therapy. The genes are
 isolated from a range of human tissues disclosed in the specification.
 The nucleic acids, proteins, antibodies and (ant)agonists are useful
 in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast
 and ovarian cancer and other cancers of the adrenal gland, bone, bone
 marrow, breast, gastrointestinal tract, liver, lung, or urogenital;
 (b) immune disorders e.g. Addison's disease, allergies, autoimmune
 haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
 disease, multiple sclerosis, rheumatoid arthritis and ulcerative
 colitis; (c) cardiovascular disorders such as myocardial ischaemia;
 (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and
 epilepsy; and (f) infectious diseases such as viral, bacterial, fungal
 and parasitic infections.
 Note: The sequence data for this patent did not form part of the
 claimed specification, but was obtained in electronic format directly
 from WFO at http://wipo.int/pub/published_pat_sequences.
 Sequence 19616 BP: 4181 A; 5695 C; 5539 G; 4201 T; 0 other;

Query Match 18.8%; Score 188.6; DB 22; Length 19616;
 Best Local Similarity 86.0%; Pred. No. 3.4e-25;
 Matches 209; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

3 TATAAAATGCATCTTTAGGCTGGGCAAGTGGCTCAGCTGTGTAATCCAGCACTTG 42
 17028 TGTGTAATAAGCTAAATGGGGCAAGCGAGTGGCTCAGCTGTGTAATCCAGCACTTG 16969
 63 GGAGGCAAGCTGGGGGATCAGCTAGGTTAGGAGTTAGAGCAACCTGGCCATCATG 122
 16968 GGAGGCAAGCTGGGGGATCAGCTAGGTTAGGAGTTAGAGCAACCTGGCCATCATG 16909
 123 GCGAAACCTGTCTCTACTAAAAATACAAAATTTAGCTGGGATGAGGCACTGTGCTCT 182
 16906 ACGAAACCTGTCTCTACTAAAAATACAAAATTTAGCTGGGATGAGGCACTGTGCTCT 16849
 183 AATCCAGCTACTCGGGAGGCTGAGACAGAAATTTGTAACCTTGAGGGGGAGGTT 242
 16848 AATCCAGCTACTCGGGAGGCTGAGGAGGAGTAATCGTTGAACCTGGGAGGAGGTT 16789
 243 GCA 245
 16788 GCA 16786


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PR 17-NOV-2000; 2000US-0249299.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250160.
PR 01-DEC-2000; 2000US-0250191.
PR 05-DEC-2000; 2000US-0251030.
PR 05-DEC-2000; 2000US-0251988.
PR 05-DEC-2000; 2000US-0256719.
PR 06-DEC-2000; 2000US-0251479.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 11-DEC-2000; 2000US-0251990.
PR 05-JAN-2001; 2000US-3254097.
PR 05-JAN-2001; 2001US-3259678.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Barash SC, Ruben SV;
XX WPI; 2001-451937/48.
XX
XX Isolated polypeptide for treating, preventing and/or prognosing
XX disorders related to the musculoskeletal system including
XX musculoskeletal cancers and also for testing and detection e.g.
XX diagnosis.
XX
XX Example 2; SEQ ID NO 3220; 751pp * Sequence Listing; English.
XX
XX The invention relates to novel genes (AAL34669; AAL37666 and proteins
XX 'AB03087-AB04109) associated with the musculoskeletal system useful
XX for preventing, treating or ameliorating medical conditions e.g. by
XX protein or gene therapy. The genes are isolated from a range of human
XX tissues disclosed in the specification. The nucleic acids, proteins,
XX antibodies and antagonists are useful in the diagnosis, treatment
XX and prevention of: (a) cancer, e.g. breast and ovarian cancer and
XX other cancers of the adrenal gland, bone, bone marrow, breast,
XX gastrointestinal tract, liver, lung, or urogenital; (b) immune
XX disorders e.g. Addison's disease, allergies, autoimmune haemolytic
XX anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's disease,
XX multiple sclerosis, rheumatoid arthritis and ulcerative colitis;
XX (c) cardiovascular disorders such as myocardial ischaemia; (d) wound
XX healing; (e) neurological diseases e.g. cerebral ataxia and epilepsy;
XX and (f) infectious diseases such as viral, bacterial, fungal and
XX parasitic infections.
XX
XX Note: The sequence data for this patent did not form part of the
XX printed specification, but was obtained in electronic format directly
XX from WIPO at ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 19616 BP; 4181 A; 5695 C; 5539 G; 4201 T; 0 other;
XX
XX Query Match 18.8%; Score 186.6; DB 22; Length 19616;
XX Best Local Similarity 86.0%; Pred. No. 3.4e-25;
XX Matches 209; Conservative 0; Mismatches 34; Indels 0; Gaps 0;
XX
XX QY 3 TATAAAATGCATCTTTAGGCTGGCCAAAGTGGCTCAGCTGTGTAATCCAGCATTG 62
XX DB 17028 TGTGTAATGCTAAATGGGCCAAGCGAGTGGCTCAGCTGTGTAATCCAGCATTG 16969
XX
XX QY 63 GGAGGCCAAGTGGCGGATACCTGAGCTTAGAGTTCAGGACCACTGGCCATCATG 122
XX DB 16969 GGAGGCCAAGTGGCGGATACCTGAGCTTAGAGTTCAGGACCACTGGCCACATG 16909
XX
XX QY 123 GCGAAACCTGTCTCTACTATAAATACAAAATTAGCTGGCATGGAGGATGTGCTGT 182
XX DB 16308 ACGAAACCTGTCTCTACTATAAATACAAAATTAGCTGGCATGGAGGATGTGCTGT 16849
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XX QY 183 AATCCAGCTACTCGGAGGCTGAGACAGAGAAATGCTTCAACTGGAGGGGAGGTT 242
XX DB 16848 AATCCAGCTACTCGGAGGCTGAGACAGAGAAATGCTTCAACTGGAGGGGAGGTT 16799
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XX QY 243 GCA 245
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Db 16788 GCA 16786

Search completed: October 24, 2003, 18:47:25
Job time : 255.249 secs


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; NAME/KEY: exon 2
; LOCATION: 2115..2262
; FEATURE:
; NAME/KEY: exon 3
; LOCATION: 3056..3202
; FEATURE:
; NAME/KEY: exon 4
; LOCATION: 3387..3472
; FEATURE:
; NAME/KEY: exon 5
; LOCATION: 5667..5923
; FEATURE:
; NAME/KEY: exon 6
; LOCATION: 7745..8955
; US-09-077-3548-3

Query Match      19.0%; Score 189.8; DB 3; Length 10390;
Best Local Similarity 86.7%; Pred. No. 1.5e-3;
Matches 209; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

QY 5 TAAAAATGCATACCTTAGGCTGGGCAAGTGGCTCAGCTCTGTATATCCAGCAGCTTTGGG 64
Db 3540 TAAAAATCTTAAGCTCTGGGCGGGCGGCGAGTGGCTCAGGCTGTATATCCAGCAGCTTTGGG 3539

QY 65 AGGCGAGGTGGGCGGATCAGCTGAGGTAGAGTTTCAGGACCACTTGGGCTATGGC 124
Db 3600 AGGCGAGGTGGGCGGATCAGCTGAGGTAGAGTTTCAGGACCACTTGGGCTATGGC 3659

QY 125 GAAACCTCTCTCTACTTAAATAACAAAAATTAGCTGGGTCATGGAGGATGTGCTGTAA 184
Db 3660 GAAACCTCTCTCTACTTAAATAACAAAAATTAGCTGGGTCATGGGCTGTAA 3719

QY 185 TCCAGCTACTCGGAGGCTGAGCAGAGAAATTGCTTGAACCTTGGAGGGGAGGTTC 244
Db 3720 TCCAGCTACTAGGAGGCTGAGCAGAGAAATTGCTTGAACCTTGGAGGGGAGGTTC 3779

QY 245 A 245
Db 3780 A 3780

RESULT 2
US-09-791-211-10
; Sequence 10, Application: US/09791211
; Patent No. 644880
; GENERAL INFORMATION:
; APPLICANT: Donna T. Ward
; APPLICANT: Andrew T. Watt
; TITLE OF INVENTION: ANTISENSE MODULATION OF WRN EXPRESSION
; FILE REFERENCE: RTS-2205
; CURRENT APPLICATION NUMBER: US/09/791,211
; CURRENT FILING DATE: 2001-02-23
; NUMBER OF SEQ ID NOS: 90
; SEQ ID NO 10
; LENGTH: 98844
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: unsure
; LOCATION: 24962
; OTHER INFORMATION: unknown
; NAME/KEY: unsure
; LOCATION: 64383
; OTHER INFORMATION: unknown
; NAME/KEY: unsure
; LOCATION: 65468
; OTHER INFORMATION: unknown
; NAME/KEY: unsure
; LOCATION: 65469
; OTHER INFORMATION: unknown
; NAME/KEY: unsure
; LOCATION: 65470
; OTHER INFORMATION: unknown

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; NAME/KEY: unsure
; LOCATION: 65471
; OTHER INFORMATION: unknown
; NAME/KEY: unsure
; LOCATION: 87130
; OTHER INFORMATION: unknown
; NAME/KEY: unsure
; LOCATION: 89049
; OTHER INFORMATION: unknown
; US-09-791-211-10

Query Match      18.9%; Score 188.8; DB 4; Length: 98844;
Best Local Similarity 86.7%; Pred. No. 3.8e-3;
Matches 208; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

QY 6 AAAAAATGCATACCTTAGGCTGGGCAAGTGGCTCAGCTCTGTATATCCAGCAGCTTTGGGA 65
Db 82007 AAAAAATGCATACCTTAGGCTGGGCAAGTGGCTCAGCTCTGTATATCCAGCAGCTTTGGGA 82066

QY 66 GGCCAAAGTGGGCGGATCAGCTGAGTTAGGAGTTTCAGACCAACTGGGCTATGGGCG 125
Db 82067 GGCCAAAGTGGGCGGATCAGCTGAGTTAGGAGTTTCAGACCAACTGGGCTATGGGCG 82126

QY 126 AACCCTGCTCTACTTAAATAACAAAAATTAGCTGGGTCATGGAGGATGTGCTGTAA 185
Db 82127 AACCCTGCTCTACTTAAATAACAAAAATTAGCTGGGTCATGGGTCATGGGTCAT 82186

QY 186 CCAGCTACTTCGGGAGGCTGAGCAGAGAAATTGCTTGAACCTTGGAGGGGAGGTTCGA 245
Db 82187 CCAGCTACTTCGGGAGGCTGAGCAGAGAAATTGCTTGAACCTTGGAGGGGAGGTTCGA 82246

RESULT 3
US-09-830-960-3/c
; Sequence 3, Application: US/09803960
; Patent No. 6187677
; GENERAL INFORMATION:
; APPLICANT: YE, Jane et al.
; TITLE OF INVENTION: ISOLATED HUMAN KINASE PROTEINS, NUCLEIC
; TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN KINASE PROTEINS, AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CL001158
; CURRENT APPLICATION NUMBER: US/09/800,960
; CURRENT FILING DATE: 2001-03-08
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 62804
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(62804)
; OTHER INFORMATION: n = A,T,C or G
; US-09-800-960-3

Query Match      18.8%; Score 185.6; DB 4; Length: 62804;
Best Local Similarity 89.3%; Pred. No. 2.7e-30;
Matches 200; Conservative 0; Mismatches 24; Indels 0; Gaps 0;

QY 22 GGCTGGGCAAGTGGCTCAGCTCTGTATATCCAGCAGCTTTGGGAGGCCAAGGTGGCGGA 81
Db 48950 GGGTGGGCAAGTGGCTCAGCTCTGTATATCCAGCAGCTTTGGGAGGCCAAGGTGGGA 48891

QY 82 TCACCTGAGGTTAGGAGTTTCAGACCAACTGGCCATCATGGCGAAACCTGTCTCTACT 141
Db 48990 TCACCTGAGGTTAGGAGTTTCAGACCAACTGGCCATCATGGCGAAACCTGTCTCTACT 48831

QY 142 AAAAAATCAAAAAATTAGCTGGGTCATGGAGGATGTGCTGTATATCCAGCTACTCGGGAG 201
Db 48830 AAAAAATCAAAAAATTAGCTGGGTCATGGAGGATGTGCTGTATATCCAGCTACTCGGGAG 48771

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QY 202 GCTGAGCAGAGAAATTCCTGTAACCTTGGAGCGGAGGTTGCA 245
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DB 48770 GCTGAGCAGAGAAATTCCTGTAACCTTGGAGCGGAGGTTGCA 48727

RESULT 4

US-579-445-26

Sequence 26, Application US/08579445

Patent No. 556653

GENERAL INFORMATION:

APPLICANT: Peruchio, Manuel

APPLICANT: Peinado, Miguel A.

APPLICANT: Iosov, Yuri

APPLICANT: Malkhsyan, Sergei

TITLE OF INVENTION: Identification of Neoplasms by Detection

TITLE OF INVENTION: of Genetic Deletions

NUMBER OF SEQUENCES: 27

CORRESPONDENCE ADDRESS:

ADDRESSEE: Knobbe, Martens, Olson & Bear

STREET: 620 Newport Center Drive, Sixteenth Floor

CITY: Newport Beach

STATE: CA

COUNTRY: U.S.A.

ZIP: 92662

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent In Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/579,445

FILING DATE:

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/08/152,484

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Kirkpatrick, Anita M.

REGISTRATION NUMBER: 32,617

REFERENCE/DOCKET NUMBER: STRATAG.009A

TELECOMMUNICATION INFORMATION:

TELEPHONE: (619) 235-8550

TELEFAX: (619) 235-0176

INFORMATION FOR SEQ ID NO: 26:

SEQUENCE CHARACTERISTICS:

LENGTH: 283 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

HYPOHETICAL: NO

ANTI-SENSE: NO

US-08-579-445-26

Query Match 18.4%, Score 184, DB 4, Length 283,

Best Local Similarity 88.8%, Pred. No. 13e-30,

Matches 199, Conservative 0, Mismatches 25, Indels 0, Gaps 0:

QY 22 GGCTGGGCAAGTGGCTCAGCTTGTAATCCAGACACTTTGGAGCGGAGGTTGCA 245

DB 1 GGCTGGGCGTGGTGGCTCACACCTGTAATCCAGACACTTTGGAGCGGAGGTTGCA 40

QY 82 TCACCTGAGTTAGGAGTTGAGAGTTCAGGACCAACCTGGCCATCATGCGGAACCTGTCTCTACT 141

DB 61 TCACCTGAGTCCAGGAGTTCAAGACCAACCTGGCCATCATGCGGAACCTGTCTCTACT 120

QY 142 AAAAAACACAAAATTAAGTGGGCGATGGAGGCGATGCTGTATATCCAGGTAATCCGGAG 201

DB 121 AAAAAACACAAAATTAAGTGGGCGGCGTGGTGGCGGCGCTGTATATCCAGGTAATCCGGAG 180

QY 202 GCTGAGCAGAGAAATTCCTGTAACCTTGGAGCGGAGGTTGCA 245

DB 181 GCTGAGCGCAGAGAAATTCCTGTAACCTTGGAGCGGAGGTTGCA 224

RESULT 5

US-58-133-629-8

Sequence 8, Application US/58133629

Patent No. 5597694

GENERAL INFORMATION:

APPLICANT: Mufree, David J.

APPLICANT: Houston, David E.

TITLE OF INVENTION: APPLICATION OF NUCLEIC ACIDS

NUMBER OF SEQUENCES: 8

CORRESPONDENCE ADDRESS:

ADDRESSEE: Wolf, Greenfield & Sacks, P.C.

STREET: 600 Atlantic Avenue

CITY: Boston

STATE: Massachusetts

COUNTRY: United States of America

ZIP: 02210

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent In Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/133,629

FILING DATE: 07-OCT-1993

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Greer, Helen

REGISTRATION NUMBER: 36,816

REFERENCE/DOCKET NUMBER: M0828/7001

TELECOMMUNICATION INFORMATION:

TELEPHONE: 617-720-3500

TELEFAX: 617-720-2441

INFORMATION FOR SEQ ID NO: 8:

SEQUENCE CHARACTERISTICS:

LENGTH: 282 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

US-08-133-629-8

Query Match 18.3%, Score 183, DB 1, Length 282,

Best Local Similarity 88.4%, Pred. No. 1.6e-30,

Matches 199, Conservative 1, Mismatches 25, Indels 0, Gaps 0:

QY 22 GGCTGGGCAAGTGGCTCAGCTTGTAATCCAGACACTTTGGAGCGGAGGTTGCA 81

DB 1 GGCTGGGCGTGGTGGCTCACACCTGTAATCCAGACACTTTGGAGCGGAGGTTGCA 60

QY 92 TCACCTGAGTTAGGAGTTGAGAGTTCAGGACCAACCTGGCCATCATGCGGAACCTGTCTCTACT 141

DB 61 TCACCTGAGTCCAGGAGTTCAAGACCAACCTGGCCATCATGCGGAACCTGTCTCTACT 120

QY 142 AAAAAACACAAAATTAAGTGGGCGATGGAGGCGATGCTGTATATCCAGGTAATCCGGAG 201

DB 121 AAAAAACACAAAATTAAGTGGGCGGCGTGGTGGCGGCGCTGTATATCCAGGTAATCCGGAG 180

QY 202 GCTGAGCAGAGAAATTCCTGTAACCTTGGAGCGGAGGTTGCA 245

DB 181 GCTGAGCGCAGAGAAATTCCTGTAACCTTGGAGCGGAGGTTGCA 224

RESULT 6

US-539-3330-37

Sequence 37, Application US/5953933D

Patent No. 6476208

GENERAL INFORMATION:

APPLICANT: Cohen, Daniel

APPLICANT: Blumenfeld, Marta

APPLICANT: Chumakov, Ilva

APPLICANT: Bougueleret, Lydie

```

APPLICANT: Bihain, Bernard
TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENES, PROTEINS AND BIALLLELIC MARKERS
FILE REFERENCE: GENSET 047AUS
CURRENT APPLICATION NUMBER: US 09/539,333D
PRIOR FILING DATE: 2000-03-30
PRIOR APPLICATION NUMBER: US 60/126,903
PRIOR FILING DATE: 1999-03-30
PRIOR APPLICATION NUMBER: US 60/131,971
PRIOR FILING DATE: 1999-04-30
PRIOR APPLICATION NUMBER: US 60/132,065
PRIOR FILING DATE: 1999-04-30
PRIOR APPLICATION NUMBER: US 60/143,928
PRIOR FILING DATE: 1999-07-14
PRIOR APPLICATION NUMBER: US 60/145,915
PRIOR FILING DATE: 1999-07-27
PRIOR APPLICATION NUMBER: US 60/146,453
PRIOR FILING DATE: 1999-07-29
PRIOR APPLICATION NUMBER: US 60/162,288
PRIOR FILING DATE: 1999-10-28
PRIOR APPLICATION NUMBER: US 09/416,384
NUMBER OF SEQ ID NOS: 231
SOFTWARE: Patent..pm
SEQ ID NO 37
LENGTH: 1154
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 1..719
FEATURE:
NAME/KEY: CDS
LOCATION: 720..1116
FEATURE:
NAME/KEY: 3'UTR
LOCATION: 1119..1154
FEATURE:
NAME/KEY: polyA signal
LOCATION: 1131..1136
FEATURE:
NAME/KEY: allele
LOCATION: 19:
OTHER INFORMATION: 8-121-197 : polymorphic base A or C
FEATURE:
NAME/KEY: allele
LOCATION: 313
OTHER INFORMATION: 8-122-271 : deletion of CAAA
FEATURE:
NAME/KEY: allele
LOCATION: 314
OTHER INFORMATION: 8-122-272 : polymorphic base A or G
FEATURE:
NAME/KEY: allele
LOCATION: 368
OTHER INFORMATION: 8-122-326 : polymorphic base A or C
FEATURE:
NAME/KEY: allele
LOCATION: 390
OTHER INFORMATION: 8-123-55 : polymorphic base A or T
FEATURE:
NAME/KEY: allele
LOCATION: 814
OTHER INFORMATION: 8-128-61 : polymorphic base G or C
FEATURE:
NAME/KEY: allele
LOCATION: 821
OTHER INFORMATION: 8-128-69 : polymorphic base C or T
FEATURE:
NAME/KEY: allele
LOCATION: 822

```

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OTHER INFORMATION: 8-128-69 : polymorphic base A or G
FEATURE:
NAME/KEY: allele
LOCATION: 838
OTHER INFORMATION: 8-128-85 : polymorphic base A or C
FEATURE:
NAME/KEY: allele
LOCATION: 897
OTHER INFORMATION: 8-129-50 : polymorphic base C or T
FEATURE:
NAME/KEY: allele
LOCATION: 908
OTHER INFORMATION: 8-129-60 : deletion of A
US-09-539-333D-37

Query Watch      : 18.3%; Score 193.6; DB 4; Length 1154;
Best Local Similarity 85.7%; Pred. No. 2.1e-30;
Matches 204; Conservative 0; Mismatches 34; Indels 3; Gaps 0;

QY 9 AAATGCATACCTTAGGCTGGGCAAGTGGCTCAAGTCTGTAAATCCAGCACTTTGGGAGG 67
D6 503 AAGAGGAAAGTCTTGGCCAGGCTCAGTGGCTCACACCTGTAAATCCAGCACTTTGGGAGG 562
QY 66 CCAAGGTGGCGGATCAGCTGAGGTAGGAGTTGAGGAGCAACCTGGCCATCGTGGAA 127
D6 563 CCGAGGCGGGAGATCAGCTGAGGTGAGGAGTTCAAGACCAAGCTTGACCAATAAGTGAA 622
QY 128 ACCCTGTCTTACTAAAAATACAAAATTTAGCTGGCATGGAGGCAATGTGCTGTAAATCC 187
D6 623 ACCCATCTCTACTAAAAATACAAAATTTAGCCAGGCAATGTGGCAGTGGCTGTAGTCC 682
QY 186 CAGTACTCGGAGGCTGAGACAGAGAAATTTGCTTGAACCTTGAGGGGAGGAGTTGCA 245
D6 583 CAGTACTCGGAGGCTGAGACAGAGAAATTTGCTTGAACCTTGAGGGGAGGAGTTGCA 743

RESULT 7
US-09-539-333D-36
; Sequence 36, Application US/0953933D
; Patent No. 6476208
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; APPLICANT: Sougouleret, Lydie
; APPLICANT: Bihain, Bernard
; APPLICANT: Essioux, Laurent
; TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENES, PROTEINS AND BIALLLELIC MARKERS
; FILE REFERENCE: GENSET 047AUS
; CURRENT APPLICATION NUMBER: US 09/539,333D
; CURRENT FILING DATE: 2000-03-30
; PRIOR APPLICATION NUMBER: US 60/126,903
; PRIOR FILING DATE: 1999-03-30
; PRIOR APPLICATION NUMBER: US 60/131,971
; PRIOR FILING DATE: 1999-04-30
; PRIOR APPLICATION NUMBER: US 60/132,065
; PRIOR FILING DATE: 1999-04-30
; PRIOR APPLICATION NUMBER: US 60/143,928
; PRIOR FILING DATE: 1999-07-14
; PRIOR APPLICATION NUMBER: US 60/145,915
; PRIOR FILING DATE: 1999-07-27
; PRIOR APPLICATION NUMBER: US 60/146,453
; PRIOR FILING DATE: 1999-07-29
; PRIOR APPLICATION NUMBER: US 60/146,452
; PRIOR FILING DATE: 1999-07-29
; PRIOR APPLICATION NUMBER: US 60/162,288
; PRIOR FILING DATE: 1999-10-28
; PRIOR APPLICATION NUMBER: US 09/416,384
; PRIOR FILING DATE: 1999-10-12
; NUMBER OF SEQ ID NOS: 231
; SOFTWARE: Patent..pm
; SEQ ID NO 36
; LENGTH: 1301

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; TYPE: DNA
; ORGANISM: Homo sapiens
; NAME/KEY: 5'UTR
; LOCATION: 1..899
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 900..1265
; FEATURE:
; NAME/KEY: 3'UTR
; LOCATION: 1266..1331
; FEATURE:
; NAME/KEY: polyA signal
; LOCATION: 1277..1282
; FEATURE:
; NAME/KEY: allele
; LOCATION: 191
; OTHER INFORMATION: 8-121-187 : polymorphic base A or G
; FEATURE:
; NAME/KEY: allele
; LOCATION: 313
; OTHER INFORMATION: 8-122-271 : deletion of CAAA
; FEATURE:
; NAME/KEY: allele
; LOCATION: 314
; OTHER INFORMATION: 8-122-272 : polymorphic base A or G
; FEATURE:
; NAME/KEY: allele
; LOCATION: 368
; OTHER INFORMATION: 8-122-326 : polymorphic base A or G
; FEATURE:
; NAME/KEY: allele
; LOCATION: 390
; OTHER INFORMATION: 8-123-55 : polymorphic base A or T
; FEATURE:
; NAME/KEY: allele
; LOCATION: 836
; OTHER INFORMATION: 8-127-25 : polymorphic base A or G
; FEATURE:
; NAME/KEY: allele
; LOCATION: 897
; OTHER INFORMATION: 8-127-119 : polymorphic base A or G
; FEATURE:
; NAME/KEY: allele
; LOCATION: 937
; OTHER INFORMATION: 8-127-159 : polymorphic base A or C
; FEATURE:
; NAME/KEY: allele
; LOCATION: 961
; OTHER INFORMATION: 8-128-61 : polymorphic base G or C
; FEATURE:
; NAME/KEY: allele
; LOCATION: 988
; OTHER INFORMATION: 8-128-68 : polymorphic base C or T
; FEATURE:
; NAME/KEY: allele
; LOCATION: 959
; OTHER INFORMATION: 8-128-69 : polymorphic base A or G
; FEATURE:
; NAME/KEY: allele
; LOCATION: 985
; OTHER INFORMATION: 8-128-85 : polymorphic base A or C
; FEATURE:
; NAME/KEY: allele
; LOCATION: 1044
; OTHER INFORMATION: 8-129-50 : polymorphic base C or T
; FEATURE:
; NAME/KEY: allele
; LOCATION: 1055
; OTHER INFORMATION: 8-129-60 : deletion of A
;
; US-09-539-333D-36
; Query Match 18.3% Score 183.6; DB 4; Length 1381;

Best Local Similarity 85.7%; Pred. No. 2,1e-30;
Matches 234; Conservative 3; Mismatches 34; Indels 3; Gaps 0;

CY 8 AAATGCATACCTTTAGGCTGGGCAAGTGGCTCAGCTGTGTAAATCCAGACATTTGGGAGG 67
|||
DB 503 AAGAGGAAGTCTTGGCCAGGCTCAGTGGCTCACACCTGTATCCAGCACATTTGGGAGG 562
|||
CY 68 CCAAGGTGGGGATCACCTGAGTTAGGATTGAGGATTCAGGACCACTTGGCATCATGCGAA 127
|||
DB 563 CCGAGGGCGGAGATCACCTGAGTTCAGGASTTCAGACCACTGACCAATATGTTGAA 622
|||
CY 128 ACCCTGCTCTACTTAAATAAGAAAATAGCTGGGCATGAGGATGTGCTGTGTAATCC 187
|||
DB 623 ACCCATCTCTACTTAAATAAGAAAATAGCCAGCATGTGGCAGTGTCTGTATGTC 682
|||
CY 198 CAGCTACTCGGAGGCTGAGACAGAGAATTCCTTGAACCTTGGAGGGGGAGGTTGCA 245
|||
DB 643 CAGCTACTCGGAGGCTGAGACAGAGAATTCCTTGAACCTGAGGAGGCGGAGGTTGCA 740

RESULT 3
US-09-539-333D-40
; Sequence 40: Application US/09539333D
; Patent No. 6476208
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; APPLICANT: Bougueleret, Lydie
; APPLICANT: Bihain, Bernard
; APPLICANT: Essieux, Laurent
; TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENES, PROTEINS AND RIBOSOMAL MARKERS
; FILE REFERENCE: GENSET/047AJS
; CURRENT APPLICATION NUMBER: US/09/539,333D
; CURRENT FILING DATE: 2000-03-30
; PRIOR APPLICATION NUMBER: US 60/126,903
; PRIOR FILING DATE: 1999-03-30
; PRIOR APPLICATION NUMBER: US 60/131,971
; PRIOR FILING DATE: 1999-04-30
; PRIOR APPLICATION NUMBER: US 60/132,065
; PRIOR FILING DATE: 1999-04-30
; PRIOR APPLICATION NUMBER: US 60/143,928
; PRIOR FILING DATE: 1999-07-14
; PRIOR APPLICATION NUMBER: US 60/145,915
; PRIOR FILING DATE: 1999-07-27
; PRIOR APPLICATION NUMBER: US 60/146,453
; PRIOR FILING DATE: 1999-07-29
; PRIOR APPLICATION NUMBER: US 60/146,452
; PRIOR FILING DATE: 1999-07-29
; PRIOR APPLICATION NUMBER: US 60/162,285
; PRIOR FILING DATE: 1999-10-28
; PRIOR APPLICATION NUMBER: US 09/416,164
; PRIOR FILING DATE: 1999-10-12
; NUMBER OF SEQ ID NOS: 231
; SOFTWARE: Patent.p
; SEQ ID NO 40
; LENGTH: 1386
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..984
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 985..1350
; FEATURE:
; NAME/KEY: 3'UTR
; LOCATION: 1351..1386
; FEATURE:
; NAME/KEY: polyA signal
; LOCATION: 1362..1367
; FEATURE:
; NAME/KEY: allele
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LOCATION: 191
OTHER INFORMATION: 8-121-187 : polymorphic base A or C
FEATURE:
NAME/KEY: allele
LOCATION: 398
FEATURE:
OTHER INFORMATION: 8-122-271 : deletion of CAAA
NAME/KEY: allele
LOCATION: 399
FEATURE:
OTHER INFORMATION: 8-122-272 : polymorphic base A or G
NAME/KEY: allele
LOCATION: 453
FEATURE:
OTHER INFORMATION: 8-122-326 : polymorphic base A or C
NAME/KEY: allele
LOCATION: 475
FEATURE:
OTHER INFORMATION: 8-123-55 : polymorphic base A or T
NAME/KEY: allele
LOCATION: 891
FEATURE:
OTHER INFORMATION: 8-127-28 : polymorphic base A or G
NAME/KEY: allele
LOCATION: 982
FEATURE:
OTHER INFORMATION: 8-127-119 : polymorphic base A or G
NAME/KEY: allele
LOCATION: 1022
FEATURE:
OTHER INFORMATION: 8-127-159 : polymorphic base A or C
NAME/KEY: allele
LOCATION: 1046
FEATURE:
OTHER INFORMATION: 8-128-6 : polymorphic base G or C
NAME/KEY: allele
LOCATION: 1053
FEATURE:
OTHER INFORMATION: 8-128-69 : polymorphic base C or T
NAME/KEY: allele
LOCATION: 1054
FEATURE:
OTHER INFORMATION: 8-128-69 : polymorphic base A or G
NAME/KEY: allele
LOCATION: 1070
FEATURE:
OTHER INFORMATION: 8-128-85 : polymorphic base A or C
NAME/KEY: allele
LOCATION: 1129
FEATURE:
OTHER INFORMATION: 8-129-50 : polymorphic base C or T
NAME/KEY: allele
LOCATION: 1140
FEATURE:
OTHER INFORMATION: 9-129-60 : deletion of A
US-09-539-333D-4C

Query Match: 18.3%; Score 183.6; DB 4, Length 1386;
Best Local Similarity 85.7%; Pred. No. 2,1e-30;
Matches 204; Conservative 0; Mismatches 34; Indels 0; Gaps 0;
QY 5 AATGCTACTTTAGCGTGGGCAAGTGGCTCAGGCTCTGTATCCAGACCTTTGGGAGG 67
DB 588 AAGAGGAAGTCTTGGCCAGGCTCAGTGGCTCACCTGTATCCAGGACCTTTGGGAGG 647
QY 68 CCAAGGTGGCGGATCAGCTTGAAGTTAGGAGTTCAGGACCAACCTGGCCATCATGGCGAA 127
DB 648 CCGAGCGGGCAGATCACCTGAGGTGAGGAGTCAAGACCAAGCTGACCAATATGTTGA 707
QY 128 ACCCTGCTCTACTAAAAATACAAAATTAGCTGGCGATGAGGAGTGTGCTGTATCC 187
DB 708 ACCCCATCTCTACTAAAAATACAAAATTAGCAGGATGTTGGCAGGTGCTGTATGCC 767
QY 188 CAGCTACTCGGAGGCTGAGACAGAGATTTCTTGAACCTTGGAGGGGAGGTTTGA 245

DB 768 CAGCTACTCGGAGGCTGAGACAGAGATTTCTTGAACCTGGAGGGGAGGTTTGA 825
RESULT 9
US-09-679-409-1
Sequence 1, Application US/09679409
Patent No. 6555316
GENERAL INFORMATION:
APPLICANT: Cohen, Daniel
APPLICANT: Blumenfeld, Marta
APPLICANT: Chumakov, Ilya
APPLICANT: Bouguieret, Lydie
APPLICANT: Essioux, Laurent
TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENE, PROTEINS AND BIALL-ELIC MARKERS
FILE REFERENCE: 53 US15.CIP
CURRENT APPLICATION NUMBER: US/09/679,409
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 09/539,333
PRIOR FILING DATE: 2000-03-03
PRIOR APPLICATION NUMBER: 09/416,384
PRIOR FILING DATE: 1999-10-12
PRIOR APPLICATION NUMBER: 60/168,088
PRIOR FILING DATE: 1999-11-30
NUMBER OF SEQ ID NOS: 134
SOFTWARE: Patent.pm
SEQ ID NO 1
LENGTH: 319608
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc feature
LOCATION: 199122..201122
OTHER INFORMATION: 5' regulatory region
NAME/KEY: exon
LOCATION: 201123..201234
OTHER INFORMATION: exon S
NAME/KEY: exon
LOCATION: 201123..201563
OTHER INFORMATION: exon S2
NAME/KEY: exon
LOCATION: 201123..201234
OTHER INFORMATION: exon T
NAME/KEY: exon
LOCATION: 215702..215746
OTHER INFORMATION: exon U
NAME/KEY: exon
LOCATION: 216836..216994
OTHER INFORMATION: exon V
NAME/KEY: exon
LOCATION: 216836..217077
OTHER INFORMATION: exon V2
NAME/KEY: exon
LOCATION: 217671..217764
OTHER INFORMATION: exon V1
NAME/KEY: exon
LOCATION: 227655..227736
OTHER INFORMATION: exon V4
NAME/KEY: exon
LOCATION: 238919
OTHER INFORMATION: exon V3
NAME/KEY: exon
LOCATION: 240440..240673
OTHER INFORMATION: exon W
NAME/KEY: exon
LOCATION: 240440..241153
OTHER INFORMATION: exon W2
NAME/KEY: exon
LOCATION: 241072..241291
OTHER INFORMATION: exon X
NAME/KEY: exon
LOCATION: 244353..244561
OTHER INFORMATION: exon Y

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? NAME/KEY: exon
? LOCATION: 246273..247802
? OTHER INFORMATION: 8-292-198 : polymorphic base A or G
? NAME/KEY: allele
? LOCATION: 207313
? OTHER INFORMATION: 8-251-322 : polymorphic base A or G
? NAME/KEY: allele
? LOCATION: 208255
? OTHER INFORMATION: 8-289-322 : polymorphic base A or G
? NAME/KEY: allele
? LOCATION: 208960
? OTHER INFORMATION: 8-287-249 : polymorphic base C or T
? NAME/KEY: allele
? LOCATION: 209123
? OTHER INFORMATION: 8-287-86 : polymorphic base A or T
? NAME/KEY: allele
? LOCATION: 209631
? OTHER INFORMATION: 8-285-319 : polymorphic base A or G
? NAME/KEY: allele
? LOCATION: 210361
? OTHER INFORMATION: 8-283-278 : polymorphic base G or C
? NAME/KEY: allele
? LOCATION: 210463
? OTHER INFORMATION: 8-283-176 : polymorphic base A or G
? NAME/KEY: allele
? LOCATION: 210486
? OTHER INFORMATION: 8-283-153 : polymorphic base G or C
? NAME/KEY: allele
? LOCATION: 210583
? OTHER INFORMATION: 8-283-56 : polymorphic base C or T
? NAME/KEY: allele
? LOCATION: 210879
? OTHER INFORMATION: 8-282-145 : polymorphic base G or C
? NAME/KEY: allele
? LOCATION: 210964
? OTHER INFORMATION: 8-282-260 : polymorphic base G or T
? NAME/KEY: allele
? LOCATION: 210979
? OTHER INFORMATION: 8-282-245 : polymorphic base A or C
? NAME/KEY: allele
? LOCATION: 211050
? OTHER INFORMATION: 8-282-174 : variable motif AAAGG or GAAGAAGGAAGGAAGGAAGG
? NAME/KEY: allele
? LOCATION: 211132
? OTHER INFORMATION: 8-282-92 : polymorphic base A or T
? NAME/KEY: allele
? LOCATION: 211247
? OTHER INFORMATION: 8-281-367 : polymorphic base A or G
? NAME/KEY: allele
? LOCATION: 211315
? OTHER INFORMATION: 8-281-299 : polymorphic base A or G
? NAME/KEY: allele
? LOCATION: 211366
? OTHER INFORMATION: 8-281-248 : polymorphic base G or C
? NAME/KEY: allele
? LOCATION: 212520
? OTHER INFORMATION: 8-279-197 : polymorphic base A or C
? NAME/KEY: allele
? LOCATION: 212821
? OTHER INFORMATION: 8-279-289 : polymorphic base C or T
? NAME/KEY: allele

Query Match      18.3%   Score 183.6; DB 4; Length 319608;
Best Local Similarity 85.7%; Pred No. 6e-30;
Matches 204; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

Cy    8  AAATGCATCTTTAGGCTGGCAAGTGGCTCAGCTGTGTAATCCACACTTTGGGAGG 67
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db    29980 AACAGGAGGCTTTGGCCAGGCTCAGTGCTCACACTGTATCCAGACTTTGGGAGG 30033
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Cy    68  CCAGGTGGCGCATCACTTAGGTTAGGTTAGGACCACTTCAGGCCATCATGGCGAA 127
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db    30040 CGAGGCGGCGCATCACTTAGGTTAGGTTAGGACCACTTCAGGCCATCATGGCGAA 30099
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

```

QY 128 ACCCTGTCTCTACTAAATAACAAAAATTAGTGGCATGGAGGCATGTGCTCTAATCC 187
Db 30100 ACCCATCTCTACTAAATAACAAAAATTAGCAGGCATGTGCGAGGTGCTCTAATCC 30159
QY 188 CAGTACTCTGGAGGCTGAGACAGAGAAATGCTTGAACCTTGAGGGGGAGGTGCA 245
Db 30160 CAGTACTCTGGAGGCTGAGACAGAGAAATGCTTGAACCTTGAGGGGGAGGTGCA 30217

RESULT 10

US-09-539-333D-1
Sequence 1, Application US/09539333D
Patent No. 6476238
GENERAL INFORMATION:
APPLICANT: Cohen, Daniel
APPLICANT: Blumenfeld, Marta
APPLICANT: Chumakov, Ilya
APPLICANT: Bougueleret, Lydie
APPLICANT: Bihain, Bernard
APPLICANT: Essioux, Laurent
TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENES, PROTEINS AND DIALELLIC MARKERS
FILE REFERENCE: GENSET 047AUS
CURRENT APPLICATION NUMBER: US/09/539,333D
CURRENT FILING DATE: 2000-03-30
PRIOR APPLICATION NUMBER: US 60/126,903
PRIOR FILING DATE: 1999-03-30
PRIOR APPLICATION NUMBER: US 60/131,971
PRIOR FILING DATE: 1999-04-30
PRIOR APPLICATION NUMBER: US 60/132,065
PRIOR FILING DATE: 1999-04-30
PRIOR APPLICATION NUMBER: US 60/143,928
PRIOR FILING DATE: 1999-07-14
PRIOR APPLICATION NUMBER: US 60/145,915
PRIOR FILING DATE: 1999-07-27
PRIOR APPLICATION NUMBER: US 60/146,453
PRIOR FILING DATE: 1999-07-29
PRIOR APPLICATION NUMBER: US 60/146,452
PRIOR FILING DATE: 1999-07-29
PRIOR APPLICATION NUMBER: US 60/152,289
PRIOR FILING DATE: 1999-10-28
PRIOR APPLICATION NUMBER: US 09/416,384
PRIOR FILING DATE: 1998-10-12
NUMBER OF SEQ ID NOS: 231
SOFTWARE: Patent.pm
SEQ ID NO 1
LENGTH: 319608
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
LOCATION: 31..1107
OTHER INFORMATION: 5' regulatory region g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 1108..1295
OTHER INFORMATION: exon A g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 14877..14920
OTHER INFORMATION: exon B g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 18778..18862
OTHER INFORMATION: exon Bbis g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 25593..25743
OTHER INFORMATION: exon C g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 29385..29502
OTHER INFORMATION: exon D g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 29367..30262
OTHER INFORMATION: exon E g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 64666..64812
OTHER INFORMATION: exon F g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 65505..65853
OTHER INFORMATION: exon G g35018 gene
FEATURE:
NAME/KEY: misc_feature
LOCATION: 65854..67854
OTHER INFORMATION: 3' regulatory region g35018 gene
FEATURE:
NAME/KEY: exon
LOCATION: 94124..94964
OTHER INFORMATION: exon g35017
FEATURE:
NAME/KEY: exon
LOCATION: 201188..201234
OTHER INFORMATION: exon S g35030 gene
FEATURE:
NAME/KEY: exon
LOCATION: 214676..214793
OTHER INFORMATION: exon T g35030 gene
FEATURE:
NAME/KEY: exon
LOCATION: 215702..215746
OTHER INFORMATION: exon U g35030 gene
FEATURE:
NAME/KEY: exon
LOCATION: 216836..216915
OTHER INFORMATION: exon V g35030 gene
FEATURE:
NAME/KEY: misc_feature
LOCATION: 213818..215618
OTHER INFORMATION: 3' regulatory region g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 215819..215941
OTHER INFORMATION: exon R complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 215819..215975
OTHER INFORMATION: exon Rbis complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 216661..216952
OTHER INFORMATION: exon Qbis complement g34572 gene
FEATURE:
NAME/KEY: exon
LOCATION: 216661..217061
OTHER INFORMATION: exon Q complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 217027..217061
OTHER INFORMATION: exon Q1 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 229647..229742
OTHER INFORMATION: exon X complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 230408..230721
OTHER INFORMATION: exon P complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 231272..231412
OTHER INFORMATION: exon Obis complement g34872 gene
FEATURE:
NAME/KEY: exon

LOCATION: 231787..231880
OTHER INFORMATION: exon 02 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 231870..231879
OTHER INFORMATION: exon 01 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 234774..234822
OTHER INFORMATION: exon 0 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 237406..237428
OTHER INFORMATION: exon Nbis complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 239719..239853
OTHER INFORMATION: exon N complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 243528..243569
OTHER INFORMATION: exon N117 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 240528..240596
OTHER INFORMATION: exon M1C90 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 240528..240617
OTHER INFORMATION: exon M1C69 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 240528..240644
OTHER INFORMATION: exon M2 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 240528..240824
OTHER INFORMATION: exon M862 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 240528..240994
OTHER INFORMATION: exon M692 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 240528..241685
OTHER INFORMATION: exon M1 complement g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 242800..242993
OTHER INFORMATION: exon M51 complement g34872 gene
FEATURE:
NAME/KEY: misc feature
LOCATION: 241686..243685
OTHER INFORMATION: 5' regulatory region g34872 gene
FEATURE:
NAME/KEY: misc feature
LOCATION: 296652..296652
OTHER INFORMATION: 3' regulatory region g34872 gene
FEATURE:
NAME/KEY: exon
LOCATION: 292653..292841

Query Match 19.34; Score 183.2; DB 4; Length 319628;
Best Local Similarity 85.34; Pred. No. 7.3e-30;
Matches 203; Conservative 1; Mismatches 34; Indels 0; Gaps 0;
Cy 8 AATGTCATCTTACGCTGGGCAAGTGGCTACGCTGTGTATCCAGACATTGGGAGG 67
Db 29990 AAGAGGAAGCTCTTGGCCAGGCTCAGTGGTCAACCTGTATCCAGACATTGGGAGG 30039

Cy 6R CCAAGTGGCGGATCACCCTGAGCTTAGGAGTTCAAGACAACTGGCCATATGCGAA 127
Db 30640 CCGAGGGGGGAGATCACCTGAGCTCAGGAGTTCAAGACCAAGTCAAGTATATGTA 30039
Cy 129 ACCCTGCTCTACTAAATACAAATAGCTGGGCGATGGAGGATGTCCTGTATCC 187
Db 30100 ACCCATCTCTACTAAATATACAAATAGCCAGGATGTGGCAGGTGCTCTATGTC 30159
Cy 198 CAGCTACTCGGAGGCTGAGACAGAAAGATTGCTTGAACCTTGGAGGGGAGGTTTCA 245
Db 30160 CAGCTACTCGGAGGCTGAGACAGGAGATTGCTTGAACCTTGGAGGGGAGGTTTCA 30217
RESULT 11
US-09-750-580-1/c
Sequence 1: Application US/09750580
Patent No. 6455280
GENERAL INFORMATION:
APPLICANT: Yen, Frances
APPLICANT: Denison, Blake
APPLICANT: Bour, Barbara
APPLICANT: Bihain, Bernard
APPLICANT: Dumas Milne Edwards, Jean-Baptiste
APPLICANT: Duclert, Aymeric
APPLICANT: Bougueleret, Lydie
APPLICANT: Edbets-Reed, Dana
APPLICANT: Salter-Cid, Luisa
TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR INHIBITING NEOPLASTIC CELL GROWTH
FILE REFERENCE: 89 US2 CIP
CURRENT APPLICATION NUMBER: US/09/750,580
CURRENT FILING DATE: 2000-12-28
PRIOR APPLICATION NUMBER: US 09/599,362
PRIOR FILING DATE: 2000-06-21
PRIOR APPLICATION NUMBER: PCT/IB99/0101
PRIOR FILING DATE: 2000-06-21
PRIOR APPLICATION NUMBER: PCT/IB99/02058
PRIOR FILING DATE: 1999-12-20
PRIOR APPLICATION NUMBER: US 49/469/099
PRIOR FILING DATE: 1999-12-21
PRIOR APPLICATION NUMBER: US 60/113,686
PRIOR FILING DATE: 1998-12-22
PRIOR APPLICATION NUMBER: US 60/441,032
PRIOR FILING DATE: 1999-06-25
NUMBER OF SEQ ID NOS: 6
SOFTWARE: Patent pro
SEQ ID NO 1
LENGTH: 81001
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc feature
LOCATION: 13946..12946
OTHER INFORMATION: 5' regulatory region
NAME/KEY: exon
LOCATION: 12947..12958
OTHER INFORMATION: exon 1
NAME/KEY: exon
LOCATION: 13470..13526
OTHER INFORMATION: exon 2
NAME/KEY: exon
LOCATION: 13641..13752
OTHER INFORMATION: exon 3
NAME/KEY: exon
LOCATION: 14271..15965
OTHER INFORMATION: exon 4
NAME/KEY: misc feature
LOCATION: 15966..17969
OTHER INFORMATION: 3' regulatory region
NAME/KEY: allele
LOCATION: 1239
OTHER INFORMATION: 20-828-311 : polymorphic base C or T
NAME/KEY: allele

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LOCATION: 12347
OTHER INFORMATION: 17-42-319 : polymorphic base C or T
NAME/KEY: allele
LOCATION: 1524
OTHER INFORMATION: 17-41-250 : polymorphic base C or T
NAME/KEY: allele
LOCATION: 42218
OTHER INFORMATION: 20-841-149 : polymorphic base A or G
NAME/KEY: allele
LOCATION: 45442
OTHER INFORMATION: 20-842-115 : polymorphic base A or G
NAME/KEY: allele
LOCATION: 77058
OTHER INFORMATION: 20-853-415 : polymorphic base C or T
NAME/KEY: primer_bind
LOCATION: 929..949
OTHER INFORMATION: 20-828.pu
NAME/KEY: primer_bind
LOCATION: 1357..1377
OTHER INFORMATION: 20-828.tp complement
NAME/KEY: primer_bind
LOCATION: 12029..12050
OTHER INFORMATION: 17-42.pu
NAME/KEY: primer_bind
LOCATION: 12581..12603
OTHER INFORMATION: 17-42.rp complement
NAME/KEY: primer_bind
LOCATION: 14992..15012
OTHER INFORMATION: 17-41.pu
NAME/KEY: primer_bind
LOCATION: 15480..15482
OTHER INFORMATION: 17-41.rp complement
NAME/KEY: primer_bind
LOCATION: 42070..42090
OTHER INFORMATION: 20-841.pu
NAME/KEY: primer_bind
LOCATION: 42572..42591
OTHER INFORMATION: 20-841.rp complement
NAME/KEY: primer_bind
LOCATION: 45329..45347
OTHER INFORMATION: 20-842.pu
NAME/KEY: primer_bind
LOCATION: 45861..45883
OTHER INFORMATION: 20-842.rp complement
NAME/KEY: primer_bind
LOCATION: 76644..76664
OTHER INFORMATION: 20-853.pu
NAME/KEY: primer_bind
LOCATION: 77166..77185
OTHER INFORMATION: 20-853.rp complement
NAME/KEY: primer_bind
LOCATION: 12201..12298
OTHER INFORMATION: 20-828-311.mis
NAME/KEY: primer_bind
LOCATION: 1240..1258
OTHER INFORMATION: 20-828-311.mis complement
NAME/KEY: primer_bind
LOCATION: 12328..12346
OTHER INFORMATION: 17-42-319.mis
NAME/KEY: primer_bind
LOCATION: 12348..12366
OTHER INFORMATION: 17-42-319.mis complement
NAME/KEY: primer_bind
LOCATION: 15222..15240
OTHER INFORMATION: 17-41-250.mis
NAME/KEY: primer_bind
LOCATION: 15242..15260
OTHER INFORMATION: 17-41-250.mis complement
NAME/KEY: primer_bind
LOCATION: 42199..42217
OTHER INFORMATION: 20-841-149.mis
NAME/KEY: primer_bind
LOCATION: 42219..42237
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OTHER INFORMATION: 20-841-149.mis complement
NAME/KEY: primer_bind
LOCATION: 45423..45441
OTHER INFORMATION: 20-842-115.mis
NAME/KEY: primer_bind
LOCATION: 45443..45461
OTHER INFORMATION: 20-842-115.mis complement
NAME/KEY: primer_bind
LOCATION: 77039..77057
OTHER INFORMATION: 20-853-415.mis
NAME/KEY: primer_bind
LOCATION: 77059..77077
OTHER INFORMATION: 20-853-415.mis complement
NAME/KEY: misc_binding
LOCATION: 1227..1251
OTHER INFORMATION: 20-828-311.probe
NAME/KEY: misc_binding
LOCATION: 12335..12359
OTHER INFORMATION: 17-42-319.probe
NAME/KEY: misc_binding
LOCATION: 42205..42230
OTHER INFORMATION: 20-841-149.probe
NAME/KEY: misc_binding
LOCATION: 45430..45454
OTHER INFORMATION: 20-842-115.probe
NAME/KEY: misc_binding
LOCATION: 77045..77075
OTHER INFORMATION: 20-853-415.probe
US-09-750-590-1

Query Match 18.2% Score 182.4; DB 4; Length 81001;
Best Local Similarity 85.0%; Pred. No. 8.3e-30;
Matches 264; Conservative 0; Mismatches 36; Indels 0; Gaps 0;

QY 6 AANAATGCATCTTTAGTGGTGGCAAGTGGCTCAGCTGTGTAAATCCAGACGACCTTTGGGA 65
DB 65052 AATAAGGAATTTGTGGCCAGTGCAGTGGCTCAGCTGTGTAAATCCAGACGACCTTTGGGA 64993
CY 66 GCGCAAGTGGCGGATCAGCTCAGGTTAGAGTTTCAGGACCAACCTGGCCATCATGCG 125
DB 64992 GCGCAAGTGGCGGATCAGCTCAGGTTAGAGTTTCAGGACCAACCTGGCCATCATGCG 64933
CY 126 AAACCTGTCTCTACTATAAAATACAAAATTAAGTGGCATGGAGCATGTGCTGTAAAT 185
DB 64932 AAACCTGTCTCTACTATAAAATACAAAATTAAGTGGCATGGAGCATGTGCTGTAAAT 64873
CY 186 CCGAGTACTCGGAGGCTGACACAGAGAGATTGTTGAACCTTGGAGGGGAGGTTGCA 245
DB 64872 CCGAGTACTCGGAGGCTGACACAGAGAGATTGTTGAACCTTGGAGGGGAGGTTGCA 64813

RESULT 12
US-09-499-522-1
; Sequence 1, Application: US/09499522
; Patent No. 6479238
; GENERAL INFORMATION:
; APPLICANT: Blumenfeld, Matta
; APPLICANT: Bouqueleret, Lydie
; APPLICANT: Bihain, Bernard
; TITLE OF INVENTION: POLYMORPHIC MARKERS OF THE LSR GENE
; FILE REFERENCE: GENSET 053AUS
; CURRENT APPLICATION NUMBER: US/09/499,522
; CURRENT FILING DATE: 2000-02-10
; EARLIER APPLICATION NUMBER: US 60/11-9,592
; EARLIER FILING DATE: 1999-02-10
; EARLIER APPLICATION NUMBER: US 60/144,784
; EARLIER FILING DATE: 1999-07-20
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: Patent.pm
; SEQ ID NO 1
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1 LENGTH: 23187
2 TYPE: DNA
3 ORGANISM: Homo sapiens
4 FEATURE:
5 NAME/KEY: exon
6 LOCATION: 2001..2356
7 OTHER INFORMATION: exon:1
8 FEATURE:
9 NAME/KEY: exon
10 LOCATION: 3543..3884
11 OTHER INFORMATION: exon:2
12 FEATURE:
13 NAME/KEY: exon
14 LOCATION: 12163..12282
15 OTHER INFORMATION: exon:3
16 FEATURE:
17 NAME/KEY: exon
18 LOCATION: 15144..15200
19 OTHER INFORMATION: exon:4
20 FEATURE:
21 NAME/KEY: exon
22 LOCATION: 15765..15911
23 OTHER INFORMATION: exon:5
24 FEATURE:
25 NAME/KEY: exon
26 LOCATION: 19579..19752
27 OTHER INFORMATION: exon:6
28 FEATURE:
29 NAME/KEY: exon
30 LOCATION: 19899..19958
31 OTHER INFORMATION: exon:7
32 FEATURE:
33 NAME/KEY: exon
34 LOCATION: 20356..20187
35 OTHER INFORMATION: exon:8
36 FEATURE:
37 NAME/KEY: exon
38 LOCATION: 20329..20957
39 OTHER INFORMATION: exon:9
40 FEATURE:
41 NAME/KEY: exon
42 LOCATION: 21047..21197
43 OTHER INFORMATION: exon:10
44 FEATURE:
45 NAME/KEY: polyA_signal
46 LOCATION: 21168..21173
47 OTHER INFORMATION: AATAAA
48 FEATURE:
49 NAME/KEY: misc_feature
50 LOCATION: 1..2000
51 OTHER INFORMATION: potential 5' regulatory region
52 FEATURE:
53 NAME/KEY: misc_feature
54 LOCATION: 22324..23187
55 OTHER INFORMATION: homology with USF2 gene in ref: e-21 Y0756;
56 FEATURE:
57 NAME/KEY: primer_bind
58 LOCATION: 523..574
59 OTHER INFORMATION: upstream amplification primer 17-2
60 FEATURE:
61 NAME/KEY: primer_bind
62 LOCATION: 1047..1068
63 OTHER INFORMATION: downstream amplification primer 17 2 , complement
64 FEATURE:
65 NAME/KEY: primer_bind
66 LOCATION: 946..963
67 OTHER INFORMATION: upstream amplification primer 99-4576
68 FEATURE:
69 NAME/KEY: primer_bind
70 LOCATION: 3385..1402
71 OTHER INFORMATION: downstream amplification primer 99-4576 , complement
72 FEATURE:
73 NAME/KEY: primer_bind
74 LOCATION: 7995..8012
75 OTHER INFORMATION: downstream amplification primer 99-4577 , complement
76
77 LOCATION: 1096..1115
78 OTHER INFORMATION: upstream amplification primer 9-19
79 FEATURE:
80 NAME/KEY: primer_bind
81 LOCATION: 1616..1615
82 OTHER INFORMATION: downstream amplification primer 9-19 , complement
83 FEATURE:
84 NAME/KEY: primer_bind
85 LOCATION: 1621..1621
86 OTHER INFORMATION: upstream amplification primer 9-20
87 FEATURE:
88 NAME/KEY: primer_bind
89 LOCATION: 2074..2093
90 OTHER INFORMATION: downstream amplification primer 9-20 , complement
91 FEATURE:
92 NAME/KEY: primer_bind
93 LOCATION: 2236..2253
94 OTHER INFORMATION: upstream amplification primer 99-4557
95 FEATURE:
96 NAME/KEY: primer_bind
97 LOCATION: 2263..2280
98 OTHER INFORMATION: downstream amplification primer 99-4557 , complement
99 FEATURE:
100 NAME/KEY: primer_bind
101 LOCATION: 2284..2102
102 OTHER INFORMATION: upstream amplification primer 9-1
103 FEATURE:
104 NAME/KEY: primer_bind
105 LOCATION: 2483..2500
106 OTHER INFORMATION: downstream amplification primer 9-1 , complement
107 FEATURE:
108 NAME/KEY: primer_bind
109 LOCATION: 2470..2489
110 OTHER INFORMATION: upstream amplification primer 9-2; , complement
111 FEATURE:
112 NAME/KEY: primer_bind
113 LOCATION: 2062..2081
114 OTHER INFORMATION: downstream amplification primer 9-2;
115 FEATURE:
116 NAME/KEY: primer_bind
117 LOCATION: 3455..3474
118 OTHER INFORMATION: upstream amplification primer 9-3
119 FEATURE:
120 NAME/KEY: primer_bind
121 LOCATION: 3882..3901
122 OTHER INFORMATION: downstream amplification primer 9-3 , complement
123 FEATURE:
124 NAME/KEY: primer_bind
125 LOCATION: 3775..3792
126 OTHER INFORMATION: upstream amplification primer 99-4558
127 FEATURE:
128 NAME/KEY: primer_bind
129 LOCATION: 4336..4356
130 OTHER INFORMATION: downstream amplification primer 99-4558 , complement
131 FEATURE:
132 NAME/KEY: primer_bind
133 LOCATION: 4302..4320
134 OTHER INFORMATION: upstream amplification primer 99-14419 , complement
135 FEATURE:
136 NAME/KEY: primer_bind
137 LOCATION: 4444..4463
138 OTHER INFORMATION: downstream amplification primer 99-14419
139 FEATURE:
140 NAME/KEY: primer_bind
141 LOCATION: 6638..6655
142 OTHER INFORMATION: upstream amplification primer 99-4577
143 FEATURE:
144 NAME/KEY: primer_bind
145 LOCATION: 7072..7093
146 OTHER INFORMATION: downstream amplification primer 99-4577 , complement
147 FEATURE:
148 NAME/KEY: primer_bind
149 LOCATION: 7995..8012
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; OTHER INFORMATION: upstream amplification primer 99-4559
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 5576..5593
; OTHER INFORMATION: downstream amplification primer 99-4559, complement
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 9622..9639
; OTHER INFORMATION: upstream amplification primer 99-3148
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 10023..10040
; OTHER INFORMATION: downstream amplification primer 99-3148, complement
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 9964..9981
; OTHER INFORMATION: upstream amplification primer 99-4560
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 10546..10563
; OTHER INFORMATION: downstream amplification primer 99-4560, complement
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 10996..11015
; OTHER INFORMATION: upstream amplification primer 99-1441, complement
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 10492..10512
; OTHER INFORMATION: downstream amplification primer 99-1441
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 11922..11930
; OTHER INFORMATION: upstream amplification primer 99-4561
; FEATURE:
; NAME/KEY: primer bind
; LOCATION: 12481..12501
; OTHER INFORMATION: downstream amplification primer 99-4561, complement

Query Match      18.2%; Score 181.8; DB 4; Length 23187;
Best Local Similarity 88.0%; Pred. No. 8.8e-30;
Matches 198; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2: AGGCTGGCCAAAGTGGCTCAGCTCTATCCAGCACTTTGGAGGCGCAGGTTGGGCGG 80
Db 5591 ACTCTGGCCACGGTGGCTCAGCTCTATCCAGCACTTTGGAGGCGCAGGTTGGGCGG 5693
QY 81 ATCACTCAGGTTAGGAGTTAGGAGTTCAGGACCAACCTGGGCATCATG37GAAACCCCTGCTTAC 140
Db 5651 ATCACTCAGGTTAGGAGTTCAGGACCAACCTGGGCATCATG37GAAACCCCTGCTTAC 5710
QY 14: TAAATAACAAATATAGTGGGCTAGGAGCATGTGCTGTATATCCAGTACTTCGGGA 200
Db 5711 TAAATAACAAAGATAGGAGGCACTGGTGGACATGCTGTATATCCAGTACTTCAGGA 5770
QY 201 GCTCAGACAGAGAAATGCTTGAACTTGGAGGGGAGGTTGCA 245
Db 5771 GCTCAGCATGAGATTCCTTGAACCCAGGAGGAGGTTGCA 5815

RESULT 13
US-09-820-002-3/c
; Sequence 3, Application US/09820022
; Patent No. 6482610
; GENERAL INFORMATION:
; APPLICANT: Gan, Weiniu
; APPLICANT: Ye, Jane
; APPLICANT: DiFrancesco, Valentina
; APPLICANT: Beasley, Ellen
; TITLE OF INVENTION: ISOLATED HUMAN PROTEASE PROTEINS.
; TITLE OF INVENTION: NUCLEIC ACID MOLECULES ENCODING HUMAN PROTEASE PROTEINS, AND
; TITLE OF INVENTION: USES THEREOF
; FILE REFERENCE: CL001194
; CURRENT APPLICATION NUMBER: US/09/820,002
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; CURRENT FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 21784
; TYPE: DNA
; ORGANISM: HUMAN
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(21784)
; OTHER INFORMATION: n = A,T,C or G
US-09-820-002-3

Query Match      19.0%; Score 180.6; DB 4; Length 21784;
Best Local Similarity 84.0%; Pred. No. 1.6e-29;
Matches 204; Conservative 0; Mismatches 39; Indels 0; Gaps 0;

QY 3 TATAAATGCATACCTTTAGGCTGGGCAAGTGGCTCAGCTCTGTAAATCCAGCAGCTTGG 62
Db 18354 TAAAAAAAACACAAATAGGCTGGGCAAGTGGCTCAGCTCTGTAAATCCAGCAGCTTGG 18295
QY 63 GSAGGCCAAAGTGGCGGATCACCTGAGGTTAGGAGTTTCAGGACCAACCTGGCCATCATG 122
Db 18294 GSAGGCCAGGCGAGGAGGATCACTTGATGTCAGGAGTTTGAGACCCAGCTGGCCACATG 18235
QY 123 GCGAAACCTGTCTCTACTTAAATAACAAAAATAGCTGGGATGGAGGATGTCGCTGT 182
Db 18234 GCGAAACCTGTCTCTACTTAAATAACAAAAATAGCTGGGATGGAGGATGTCGCTGT 18175
QY 183 ATCCCGAGCTACTCGGAGGCTGAGACAGAGAAATGCTTGAACTTGGAGGGGAGGTT 242
Db 18174 ATCCCGAGCTACTCGGAGGCTGAGACAGAGAAATGCTTGAACTTGGAGGGGAGGTT 18115
QY 243 GCA 245
Db 18114 GCA 18112

RESULT 14
US-09-443-184-35
; Sequence 35, Application US/09443184A
; Patent No. 6372431
; GENERAL INFORMATION:
; APPLICANT: Cunningham, Mary Jane
; APPLICANT: Zweiger, Gary
; APPLICANT: Kaser, Matthew R.
; APPLICANT: Panzer, Scott
; APPLICANT: Seilhammer, Jeffrey J.
; APPLICANT: Yue, Henry
; APPLICANT: Baughn, Mariah
; APPLICANT: Azimzal, Yada
; APPLICANT: Ja, Preeti
; TITLE OF INVENTION: MAXIMALIAN TOXICOLOGICAL RESPONSE MARKERS
; FILE REFERENCE: PC-0307 US
; CURRENT APPLICATION NUMBER: US/09/443,184A
; NUMBER OF SEQ ID NOS: 138
; SOFTWARE: PERL Program
; SEQ ID NO 35
; LENGTH: 1762
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. 6372431 2742442CB1
US-09-443-184-35

Query Match      19.0%; Score 180.2; DB 4; Length 1762;
Best Local Similarity 84.2%; Pred. No. 1.2e-29;
Matches 203; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 5 TAAAAATGCATACCTTTAGGCTGGGCAAGTGGCTCAGCTCTGTAAATCCAGCAGCTTGG 64
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DB 1458 TTAAGTAAATACTTTGGGCTGGCACAGTGGCTCACACCTGTATATCCCGACACTTGGG 1517
QY 65 AGGCAAGGTGGCGGATACCTTGAGGTTAGGAGTTACAGGACCAACTGGCCATCATGGC 124
DB 1518 AAGCTAGGTGGCGACATCAGTTGAGGTGAGGATCTTAGACGAGCTGGTCAACATGGC 1577
QY 125 GAAACCTGCTCTTACTAAATAATACAAANATTAGGTGGGCTATGGAGCATGTGCTGTAA 164
DB 1578 GAAACCCATCTCTTACTAAATAATACAAANATTAGGTGGGCTATGGAGCATGTGCTGTAA 1637
QY 185 TCCAGTACTCGAGAGGCTGAGACAGAGAAATGGTTGAACCTTGGAGGGGAGGCTTGC 244
DB 1638 TCCAGTACTTGGAGGCTGAGGACAGAAATGGGTTGAACCCGGGAGGCTGAGGCTTGC 1697
QY 245 A 245
DB 1698 A 1698

RESULT 15

US-09-491-356C-1/c
Sequence 1, Application US/0949135C
Patent No. 5566061
GENERAL INFORMATION:
APPLICANT: Philibert, Robert A.
APPLICANT: Ginns, Edward I.
APPLICANT: Deliss, Lynn
TITLE OF INVENTION: IDENTIFICATION OF POLYMORPHISMS IN THE PCT94 REGION OF XOL2
FILE REFERENCE: 9465-60511
CURRENT APPLICATION NUMBER: US/09/491.356C
CURRENT FILING DATE: 2000-01-26
PRIOR APPLICATION NUMBER: PCT/US99/09365
PRIOR FILING DATE: 1999-04-29
PRIOR APPLICATION NUMBER: 63/083,465
PRIOR FILING DATE: 1998-04-29
NUMBER OF SEQ ID NOS: 24
SOFTWARE: PatentIn version 3.1
SEQ ID NO 1
LENGTH: 55298
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc feature
LOCATION: (485)..(485)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (498)..(498)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (16728)..(16728)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (22750)..(22750)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (22756)..(22756)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (28519)..(28519)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (44804)..(44804)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (45002)..(45002)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (54049)..(54049)
OTHER INFORMATION: n is not determined
NAME/KEY: misc feature
LOCATION: (54226)..(54226)
OTHER INFORMATION: n is not determined

US-09-491-356C-1

Query Match 18.0%; Score 180; DB 4; Length 55298;
Best Local Similarity 83.6%; Pred. No. 2,5e-29;
Matches 204; Conservative 0; Mismatches 40; Indels 0; Gaps 0;
QY 2 ATATATAAATGGCATACTTTAGGCTGGGCAAGTGGCTCAGGTCAGTCTGTATATCCCGACACTTT 61
DB 43196 ATCTAGAAAAGCACTTAGAGGCTGGGCTGATGGCTCATGCTGTATATCCCGACACTTT 43119
QY 62 GGGAGGCCAAGGTGGGCGGATACCTGAGGTTAGGAGTTACAGGACCAACTGSCCATCAT 121
DB 43196 GGGAGGCCAAGGTGGGCGGATACCTGAGGTCAGGTCAGGAGTTACAGGACCAACTGSCCATCAT 43129
QY 122 GGGGAAACCCCTGCTCTACTATAAATAATACAAANATTAGGTGGGCTATGGAGCATGTGCTGT 191
DB 43078 GGTGAAACCCCTGCTCTACTATAAATAATACAAANATTAGGTGGGCTATGGAGCATGTGCTGT 43019
QY 192 TATCCCGACCTACTCGGGAGGCTGAGACAGAAATTTGTTGAACCTTGGAGGGGAGGT 241
DB 43019 TATCCCGACCTACTCGGGAGGCTGAGGAGGCTGAGGAGGAGGAGGAGGAGGAGGAGGAGG 42959
QY 242 TGCA 245
DB 42958 TGTA 42955

Search completed: October 24, 2003, 15:43:01

Job time : 66.6458 secs


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Dh 2643 ATCTCTACTATAAATAACAAAATAGCTGGGGCTGGTGGGGTGGCTATATATCCCAAGCT 2972
Qy 193 ACTGGGAGGCTGAGACAGAGAGATTGTTGAACTTTGGAGGGGGAGATTGCA 245
Dh 2993 ACTGGGAGGCTGAGAAAGAGAGATCGTTGAAACCCGAGGAGGAGTTGCA 2955

RESULT 2
US-09-764-891-6869
; Sequence 6869, Application US/09764891
; Publication No. US20030377808A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC006
; CURRENT APPLICATION NUMBER: US/09/764,891
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PAM or file wrapper
; NUMBER OF SEQ ID NOS: 10231
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 6869
; LENGTH: 30573
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-891-6869

Query Match 19.1%; Score 191.4; DB 11; Length 30573;
Best Local Similarity 90.7%; Pred. No. 5.1e-28;
Matches 204; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 21 AGGCTGGCAAGTGGCTCAGCTGTGTAATCCACAGATTTGGAGGCGCAGGTGGGCG 80
Dh 6984 AGGCTGGGCAAGTGGCTCAGCTGTGTAATCCACAGATTTGGAGGCGCAGGTGG 7043
Qy 8 ATCACTGAGGTTAGGAGTTGAGGATTCAGGACCAAGCTGGCCATCAAGGCAACCTGTCTCTAC 140
Dh 7044 ATCACTGAGGTCAGGAGTTGAGGATTCAGGACCAAGCTGGCCATCAAGGCAACCTGTCTCTAC 7103
Qy 141 TAAATAACAAAATAGTGGGATGAGGAGGATGTGCTGTAAATCCAGTACTCGGCA 200
Dh 7104 TAAATAACAAAATAGTGGGATGAGGAGGATGTGCTGTAAATCCAGTACTCGGCA 7163
Qy 201 GCTGAGACAGAGAGATTGCTTGAACCTTGGAGGCGGAGGTTGCA 245
Dh 7164 GCTGAGGCGAGAGAGATTGCTTGAACCTTGGAGGCGGAGGTTGCA 7203

RESULT 3
US-09-764-891-6868
; Sequence 6868, Application US/09764892
; Publication No. US20030377808A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC006
; CURRENT APPLICATION NUMBER: US/09/764,891
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PAM or file wrapper
; NUMBER OF SEQ ID NOS: 10231
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 6868
; LENGTH: 32195
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-891-6868

Query Match 19.1%; Score 191.4; DB 11; Length 32195;
Best Local Similarity 90.7%; Pred. No. 5.3e-28;
Matches 204; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 21 AGGCTGGCAAGTGGCTCAGCTGTGTAATCCACAGATTTGGAGGCGCAGGTGGGCG 80
Dh 12766 AGGCTGGGACAGTGGCTCAGCTGTGTAATCCACAGATTTGGAGGCGCAGGTGG 12925
```

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Qy 81 ATCACTGAGGTTAGGAGTTGAGGATTCAGGACCAAGCTGGCCATCATGCGAATACCCCTGTCTAC 140
Dh 12826 ATCACTGAGGTCAGGAGTTCAAGACCAAGCTGGCCCAACATGTTGAACCTGTCTCTAC 12885
Qy 141 TAAATAACAAAATAGTGGGATGAGGAGGATGTGCTGTAAATCCAGTACTCGGCA 200
Dh 12886 TAAATAACAAAATAGTGGGATGAGGAGGATGTGCTGTAAATCCAGTACTCGGCA 12945
Qy 201 GCTGAGACAGAGAGATTGCTTGAACCTTGGAGGCGGAGGTTGCA 245
Dh 12946 GCTGAGGCGAGAGAGATTGCTTGAACCTTGGAGGCGGAGGTTGCA 12990

RESULT 4
US-09-914-353-21993
; Sequence 21993, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; THERAPY OF OVARIAN CANCER
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07
; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 21993
; LENGTH: 7677
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: 1..7677
; OTHER INFORMATION: n = A,T,C or G
US-09-814-353-21993

Query Match 19.1%; Score 190.8; DB 12; Length 7677;
Best Local Similarity 60.7%; Pred. No. 3.9e-28;
Matches 168; Conservative 0; Mismatches 22; Indels 11; Gaps 3;

Qy 1 AAATAAAAATGATACCTTAGCTGGGCAAGTGGCTCAGCTGTGTAAATCCAGCACTT 60
Dh 6936 AAATAAAAATGATACCTTAGCTGGGCGGCTCAGCTGTGTAAATCCAGCACTT 6895
Qy 61 TGGAGGCGCAAGTGGGCGGATCACCCTGAGGTTAGGAGTTGAGGATTCAGGACCACTGCGCATCA 120
Dh 6896 TGGAGGCGCAAGTGGGCGGATCACCCTGAGGTTAGGAGTTGAGGATTCAGGACCACTGCGCATCA 6955
Qy 121 TGGGAAACCCCTGTCTCTACTTAAATAATACAAAATTAGCTGGGCAATGGAGGCAATGTGCCT 180
Dh 6956 TGGGAAACCCCTGTCTCTACTTAAATAATACAAAATTAGCGGGCGCTGGTGGCACTTGCCT 7015
Qy 181 GTAATCCCACTACTCGGAGGCTGAGACAGAGAAATTCCTGAACCTTGGAGGGGAGG 240
Dh 7016 GTAATCCCACTACTCGGAGGCTGAGGACAGAGAAATTCGATTAAACCTGGAGGGGAGA 7075
Qy 241 TTGC-----ATATCTGAGTGTGAAATGTGATTTCTTTTCTCTTCTGTATTTTG 296
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Db 7076 TTGCAATGAGTCGACACCTGCTCCAGCTGGGTGACACAGAGAGACACTGCTCTGG 7135
QY 297 AACTTTTCTATAAATGATGTGTTTGTGTTTATATGGAATAATATTATGCTTCAAT 356
Db 7136 AAAAAAAGAAATCTCACTCACTACTAGAGAGGATGTCAGAAATTCACGATTCAGGT 7195
QY 357 GTATATACCTAAGAACTAAGACACAGTAAATATATATAGTATAGCATTTATAGGT 416
Db 7196 CTTGAACCTTGGATATGCAAGAAGGATATATAAATATTTTCATTATGATCAGTTT 7255
QY 417 TTTCTGTGTAGGAGATCAACATAGAAATATATATTAAATGGGTGACATAATTTCTAAGA 476
Db 7256 TTAAGGCTTTGAGCT---TCTATAGTGTCTCAGATCCCACTAGATAATTTAAAGC 7312
QY 477 ATACATACAGTATATTTTATACATTA---AGAAACACACACATCACTATTAT 532
Db 7313 ATCATATTAGAAATACCTTTAGAGAGACTATATAAGAAATAGAGATTTGAAATTAC 7372
QY 533 CCATGATTTGTTAACCACATATACCTGTTGATATTTGTTATTCATGCTGCTTTTAA 592
Db 7373 AAGGATTTGTTTCAATTAAGACCCAGATCTTAAGTTTCTTTTGAATTCAGTTAA 7432
QY 593 AAATCT 598
Db 7433 ACAAT 7433
RESULT 5
US-09-836-613-3
Sequence 3, Application US/39836613
Publication No. US20030039643A1
GENERAL INFORMATION:
APPLICANT: HOPKINS, JOHN JOSEPH; SCOTT, HAMISH STEELE;
WEBER, BERGIT; BLANCH, LIANNE; ARSON, DONALD STEWART
TITLE OF INVENTION: SYNTHETIC MAXIMALIAN
N-ACETYLGLUCOSAMINIDASE AND GENETIC SEQUENCES
NUMBER OF SEQUENCES: 6
ENCODING SAME
CORRESPONDENCE ADDRESS:
ADDRESSEE: NIXON PEARCE LLP
STREET: 990 STEWART AVENUE
CITY: GARDEN CITY
STATE: NEW YORK
COUNTRY: UNITED STATES
ZIP: 11530
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.25
CURRENT APPLICATION DATA: US/03/836,613
APPLICATION NUMBER: US/03/836,613
FILING DATE: 17-Apr-2001
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US96/00747
FILING DATE: 22-NOV-1996
ATTORNEY/AGENT INFORMATION:
NAME: POZALSKY, ANN R.
REGISTRATION NUMBER: 34,697
REFERENCE/DOCKET NUMBER: 2243/104
TELECOMMUNICATION INFORMATION:
TELEPHONE: 516 742 4343
TELEFAX: 516 742 4366
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 10380 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Homo sapiens

POSITION IN GENOME:
CHROMOSOME/SEGMENT: Chromosome 17
FEATURE:
NAME/KEY: exon 1
LOCATION: 990...1372
FEATURE:
NAME/KEY: exon 2
LOCATION: 2115...2262
FEATURE:
NAME/KEY: exon 3
LOCATION: 3056...3232
FEATURE:
NAME/KEY: exon 4
LOCATION: 3387...3472
FEATURE:
NAME/KEY: exon 5
LOCATION: 5567...5923
FEATURE:
NAME/KEY: exon 6
LOCATION: 7745...8955
SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-09-836-613-3
Query Match 19.0% Score 189.8; DB 11; Length 10380;
Best Local Similarity 86.7%; Pred.No. 7e-28;
Matches 209; Conservative 0; Mismatches 32; Indels 0; Gaps 0;
QY 5 TAAAAATGCATACTTTTAGGCTGGGCAAAAGTGGCTCAGCTGTGTAAATCCGACGACTTTGGG 64
Db 3540 TAAAAATTAAGCTCTGGGCGGGCGAGTGGCTCAGCTGTGTAAATCCGACGACTTTGGG 3599
QY 65 AGGCCAAGGTGGGCGGATCACCTAGGTTAGGAGTTAGGAGTTCAGAGCAACAACTGGCCATCATGGC 124
Db 3600 AGGCCAAGGTGGGCGGATCACCTAGGTTAGGAGTTCAGAGCAACAACTGGCCATCATGGC 3659
QY 125 GAAACCTGTCTTACTTAAATAACAAAATTAAGCTGGGCTAGGAGTTCAGAGCAACAACTGGCCTGTAA 184
Db 3660 GAAACCTGTCTTACTTAAATAACAAAATTAAGCTGGGCTAGGAGTTCAGAGCAACAACTGGCCTGTAA 3719
QY 185 TCCAGCTACTCGGAGGCTGAGACAGAGAAGTAATGTTGAACCTTGAGGGGGAGGTTC 244
Db 3720 TCCAGCTACTCGGAGGCTGAGACAGAGAAGTAATGTTGAACCTTGAGGGGGAGGTTC 3779
QY 245 A 245
Db 3780 A 3780
RESULT 6
US-10-027-632-123771
Sequence 123771, Application US/10027632
GENERAL INFORMATION:
APPLICANT: Wang, David G.
TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
Polymorphisms in the Human Genome
FILE REFERENCE: 108827.129
CURRENT APPLICATION NUMBER: US/10/027,632
FILING DATE: 2002-04-30
PRIOR APPLICATION NUMBER: US 60/218,806
PRIOR FILING DATE: 2000-07-12
PRIOR APPLICATION NUMBER: US 60/198,676
PRIOR FILING DATE: 2000-04-20
PRIOR APPLICATION NUMBER: US 60/193,483
PRIOR FILING DATE: 2000-03-29
PRIOR APPLICATION NUMBER: US 60/195,218
PRIOR FILING DATE: 2000-02-24
PRIOR APPLICATION NUMBER: US 60/157,163
PRIOR FILING DATE: 1999-11-23
PRIOR APPLICATION NUMBER: US 60/156,358
PRIOR FILING DATE: 1999-09-28
PRIOR APPLICATION NUMBER: US 60/146,002
PRIOR FILING DATE: 1999-08-09
NUMBER OF SEQ ID NOS: 325723


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RESULT 9
US-10-027-632-102144/c
; Sequence 102144, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; POLYMORPHISMS IN THE HUMAN GENOME
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; PRIOR FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,359
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 102144
; LENGTH: 2490
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-102144

Query Match      18.9%; Score 189.4; DB 13; Length 2490;
Best Local Similarity 87.0%; Pred. No. 4.8e-28;
Matches 206; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

US-10-027-632-102144
QY 6 AAAAAATGTCATCTTTAGGCTGGGCAAGTGGCTCAGCTCTGTAAATCCAGCAGCATTGGGA 65
DB 476 AAAAAAAAAAAAAAGGTCGSCATGTTGCTCAGCTGTAAATCCAGCAGCATTGGGA 417
QY 66 GGGCAAGGTGGCGGATCACCCTGAGGTAGGAGTTGAGGACCAACCTGGCCATCATGCG 125
DB 416 GGGCGAGGTGAGCGGATCACCCTGAGGTAGGAGTTGAGGACCAACCTGGCCATCATGCG 357
QY 126 AAACCTGTCTTACTTAAATACAAAATTAGCTGGGATGGGATGGGATGTCCTGTAA 185
DB 356 AAACCCGCTCTTACTTAAATACAAAATTAGCTGGGATGGGATGGGATGTCCTGTAA 297
QY 186 CCCAGCTACTCGGAGGCTGAGACAGAGAAATTTGTTGAACCTTGGAGGGAGGTTGC 244
DB 296 CCCAGCTACTCGGAGGCTGAGGAGGAGAAATGGCTTGAACCTTGGAGGGAGGTTGC 238

RESULT 10
US-10-027-632-111607
; Sequence 111607, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; POLYMORPHISMS IN THE HUMAN GENOME
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; PRIOR FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 111607
; LENGTH: 2490
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-111607

Query Match      18.9%; Score 188.8; DB 13; Length 2490;
Best Local Similarity 85.4%; Pred. No. 3.7e-28;
Matches 205; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

US-10-027-632-111607
QY 14 ATACTTGGCTGGGCAAGTGGCTCAGCTCTGTAAATCCAGCAGCATTGGGAGGCAAGG 73
DB 244 ATATTAGGCTGGGCTGGCTCAGCTCTGTAAATCCAGCAGCATTGGGAGGCAAGG 303
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QY      6  AAAAAATGCATACCTTTAGGCTGGCAAGTGGCTCAGCTCTGTATNTCCAGACACTTTGGGA  65
      |||||  ||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db     13401  AAAAAATCACTCTTTGGGTGGCGCAGTGGCTCACACCTGTAAATCCAGCACCTTTGGGA  13460
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QY      66  GGCCAGGTGGCGGATCACCTGAGGTAGGAGTTCAGGACCAACCTGGCGATCATGGCG  125
      |||||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db     13461  GGCCAGGTGGCGGATCACCTGAGGTAGGAGTTCAGGACCAACCTGGCGATCATGGCG  13520
      |||||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
QY     126  AAACCCCTGTCTTACTTAAATAACAAAAATAGCTGGGCATGGAGGATGTGCTGTAAAT  185
      |||||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db     13521  AAACCCCATCTCTACTTAAANATACAAAAATAGCTGGGTGTGTTGGCACACACCTGTAAT  13580
      |||||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
QY     186  CCGAGTACTCGGAGGCTGAGACAGAGAGATTTGTAACCTTGGAGGGGAGGTTGCA  245
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Db     13581  CCGAGTACTTGGGAGGCTGAGGCAAGAGATAGCTTGAACCTGGAGATGGAGGTTGCA  13640
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RESULT 15

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US-10-091-504-1599
; Sequence 1593, Application: US/10091504
; Publication No.: US20030059908A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC007C1
; CURRENT APPLICATION NUMBER: US/10/091-504
; CURRENT FILING DATE: 2002-03-07
; NUMBER OF SEQ ID NOS: 2442
; Prior Application removed - See File Wrapper or Paim
; SOFTWARE: PatentIn Ver. 2.3
; SEQ ID NO 1599
; LENGTH: 17792
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-10-091-504-1599
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Query Match      18.9%; Score 188.8; DB 14; Length 17792;
Best Local Similarity 86.7%; Pred. No. 1.4e-27;
Matches 209; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

QY      6  AAAAAATGCATACCTTTAGGCTGGCAAGTGGCTCAGCTCTGTAAATCCAGCACCTTTGGGA  65
      |||||  ||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db     13401  AAAAAATCACTCTTTGGGTGGCGCAGTGGCTCACACCTGTAAATCCAGCACCTTTGGGA  13460
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QY      66  GGCCAGGTGGCGGATCACCTGAGGTAGGAGTTCAGGACCAACCTGGCGATCATGGCG  125
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Db     13461  GGCCAGGTGGCGGATCACCTGAGGTAGGAGTTCAGGACCAACCTGGCGATCATGGCG  13520
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QY     126  AAACCCCTGTCTTACTTAAATAACAAAAATAGCTGGGCATGGAGGATGTGCTGTAAAT  185
      |||||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db     13521  AAACCCCATCTCTACTTAAANATACAAAAATAGCTGGGTGTGTTGGCACACACCTGTAAT  13580
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QY     186  CCGAGTACTCGGAGGCTGAGACAGAGAGATTTGTTGAACCTTGGAGGGGAGGTTGCA  245
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Search completed: October 24, 2003, 18:34:54
Job time : 250.585 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - Nucleic search, using sw model

Run on: October 24, 2003, 18:11:56 Search time 2091.3 Seconds
Without alignments:
14633.309 Million cell updates/sec

Title: US-09-830-902-1_COPY_100000_101000
Perfect score: 1001
Sequence: 1 aatataaaagcatactt.....accgattcaggctcttttt 1501

Scoring table: IDENTITY_NJC

Searched: Gapop 10.0, Gapext 1.0

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 5%
Maximum Match 100%
Clustering first 45 summaries

Database:

1: em_esth1.*

2: em_esth2.*

3: em_esth3.*

4: em_esth4.*

5: em_esth5.*

6: em_esth6.*

7: em_esth7.*

8: em_esth8.*

9: gk_est1.*

10: gk_est2.*

11: gk_est3.*

12: gk_est4.*

13: gk_est5.*

14: gk_est6.*

15: em_estfun.*

16: em_estfun.*

17: em_gss_num.*

18: em_gss_inv.*

19: em_gss_pln.*

20: em_gss_vrt.*

21: em_gss_fun.*

22: em_gss_nam.*

23: em_gss_mus.*

24: em_gss_pro.*

25: em_gss_rtd.*

26: em_gss_pht.*

27: em_gss_vrl.*

28: gk_gss1.*

29: gk_gss2.*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	ID	Description
1	530.4	53.0	556	14	CB241186
2	512.2	51.2	532	9	AW37847
3	403.3	40.3	743	13	BUS53405
4	398.2	39.8	877	13	BUI93776

5	388.6	38.8	784	10	BC204578
6	388	38.8	749	10	BC217921
7	350	35.0	694	10	BC193969
8	266.2	26.6	728	10	BF216343
9	277.8	27.8	760	10	BG185583
10	244.6	24.4	460	10	BE175740
11	218.2	21.8	239	10	BG206984
12	205.4	20.5	421	14	W42916
13	190.4	19.0	516	23	AQ112451
14	169.8	16.9	515	14	CA946732
15	167.4	16.7	453	9	A1339725
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17	167.4	16.7	562	9	A1708237
18	167.2	16.7	560	28	AC057239
19	167.2	16.7	711	28	AQ415030
20	187	18.7	499	2	HSMD36977
21	166.8	16.6	363	9	AA493808
22	166.6	16.6	500	3	AW338860
23	166.6	16.6	826	13	BQ438665
24	166.4	16.6	420	2	HSMD35526
25	166.4	16.6	410	28	AC073107
26	166.4	16.6	994	12	BM679479
27	166.2	16.6	368	9	AA633264
28	166.2	16.6	397	9	A1821805
29	166.2	16.6	407	9	A1821056
30	166.2	16.6	425	9	A1922092
31	166.2	16.6	439	9	A1927119
32	166.2	16.6	442	28	AC018157
33	166.2	16.6	600	13	BC093702
34	166.2	16.6	707	28	AO540344
35	165.8	16.6	544	9	AL599310
36	165.6	16.5	442	10	BF991881
37	165.6	16.5	563	28	AQ91420
38	165.6	16.5	662	29	AG37584
39	165.4	16.5	361	9	AW65668
40	165.4	16.5	700	28	AQ117150
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44	185	18.5	395	9	AA278496
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ALIGNMENTS

RESULT:

CB241186

LOCUS

DEFINITION

CB241186

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PubMed

COMMENT

CB241186 556 bp mRNA linear EST 12-PEB-2003
UT-CP-FNC-afx-b-20-C-UI s1 UT-CP-FNC Homo sapiens cDNA clone
UT-CP-FNC-afx-b-20-C-UI 3, mRNA sequence.

CB241186.1 GI:28162930

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Euteria; Primates; Catarrhini; Homidae; Homo.

1 bases 1 to 556

Normalisation and subtraction: two approaches to facilitate gene

discovery

Genome Res. 6 (9), 791-806 (1996)

97044477

8989548

Contact: McCray, PB

McCray Lab

University of Iowa

2024 University of Iowa Med Labs, Iowa City, IA 52242, USA

Tel: 319 356 4566

Fax: 319 356 7171

E-mail: pau-mccray@iowa.edu

Tissue Procurement: Dr. M. J. Welsh, University of Iowa

cDNA library preparation: Dr. M. Bento Soares, University of Iowa
 cDNA library Arrayed by: Dr. M. Bento Soares, University of Iowa
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
 Clone Distribution: Researchers may obtain clones from Research
 Genetics (www.resgen.com) or from Open Biosystems
 (www.openbiosystems.com).
 The following repetitive elements were found in this cDNA
 sequence: 1-22, >AT-rich=Low_complexity (matched complement):
 Seq primer: M13 FORWARD
 POLYA=Yes.

FEATURES
 source
 .. 556
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="UI-CF-FNO-afx-b-20-0-UI"
 /tissue_type="Human Lung Epithelial cells"
 /lab_host="DH10B (Life Technologies) (T^r phage resistant)"
 /clone_lib="UI-CF-FNO"
 /note="Organ: Lung; Vector: pT73-Pac (Pharmacia) with a
 modified poly-linker; Site 1: EcoR I; Site 2: Not I;
 UI-CF-FNO is a subtracted cDNA library derived from two
 normalized Human lung epithelial cell libraries (EN1 and
 EN2). The library was subtracted according to according to
 Bonaldi, Lennon and Soares, Genome Research, 6:751-806,
 1996. For additional information, contact:
 Bento-soares@uiowa.edu
 TAG LIB=UI-CF-FNO
 TAG TISSUE=Lung Epithelial Cells Tissue nos 359-363
 TAG_SEQ=GCGGTAGGC"
 BASE COUNT 196 a 93 c 187 t
 ORIGIN

Query Match 53.0%; Score 530.4; DE 14; Length 556;
 Best Local Similarity 99.8%; Pred. No. 5; Seq-Id
 Matches 53%; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 391 TATATTAGTATACGATTTATTAGTTCTTTTGTAGCGATGACATAGAAATATATT 450
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 16 TATATTAGTATACGATTTATTAGTTCTTTTGTAGCGATGACATAGAAATATATT 75
 QY 451 TAAATGGCTGACATAATTTCTAGAAATACATACAGTATATTTTATACATTAAGAA 510
 Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 76 TAAATGGCTGACATAATTTCTAGAAATACATACAGTATATTTTATACATTAAGAA 135
 QY 511 ACAGGAGCATCATCTTANTCCATCATTTGTTTACACCATATACCTGTGTGATTT 570
 Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 136 ACAGGAGCATCATCTTANTCCATCATTTGTTTACACCATATACCTGTGTGATTT 195
 QY 571 TGTATTGTCATGTGCTTTTAAATCTAGATAGAGAAATATCGATTCTCTACTTCACT 630
 Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 196 TGTATTGTCATGTGCTTTTAAATCTAGATAGAGAAATATCGATTCTCTACTTCACT 255
 QY 631 GAATCTTTGAAAAAATAAACCAGCGTCAGCGCTTCAAACTTTAGAACGGTACATACCT 690
 Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 256 GAATCTTTGAAAAAATAAACCAGCGTCAGCGCTTCAAACTTTAGAACGGTACATACCT 315
 QY 691 TGGAAACAGACATTTGGAGATACACCTGTTTAAAGGAAATACCTTTGTAACCTGCAAGAC 750
 Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 316 TGGAAACAGACATTTGGAGATACACCTGTTTAAAGGAAATACCTTTGTAACCTGCAAGAC 375
 QY 751 ATTATTACTTAAAGAGAAACACAAAGATCTTCATGATGCTGTCGCTTACAGAACACAC 810
 Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 376 ATTATTACTTAAAGAGAAACACAAAGATCTTCATGATGCTGTCGCTTACAGAACACAC 435
 QY 811 CTAAGTTTACAGGACTTTTATAGTCTTACATATTTTGTSCACCAAACTTGAAGATGAAC 870
 Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 436 CTAAGTTTACAGGACTTTTATAGTCTTACATATTTTGTSCACCAAACTTGAAGATGAAC 495
 QY 871 AGAAACAGACTTAAACAAAATATACAAATGCAATGTAAATTTTCTGTTTT 922
 Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 496 AGAAACAGACTTAAACAAAATATACAAATGCAATGTAAATTTTCTGTTTT 547

RESULT 2
 AM297897 532 bp mRNA linear EST 16-JAN-2000
 LOCUS UI-H-BKO-ajv-f-11-0-JI.s1 NCI_CGAP_Sub6 Homo sapiens cDNA clone
 DEFINITION IMAGE:2733092.1, mRNA sequence.
 ACCESSION AM297897
 VERSION AM297897.1 GI:16724522
 KEYWORDS EST.
 SOURCE Homo sapiens (human);
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 532)
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-rs@mail.nih.gov
 The sequence contained an oligo-dT track that was present in the
 oligonucleotide that was used to prime the synthesis of first
 strand cDNA and therefore this may represent a bonafide poly A
 tail...cDNA library Preparation: M.B. Soares Lab Clone distribution:
 NCI-CGAP clone distribution information can be found through the
 : M.A.G.E. Consortium/LLNL at:
 www.bio.llnl.gov/bbrp/image/image.html The following repetitive
 elements were found in this cDNA sequence: 1-23,
 >AT-rich=Low_complexity
 Seq primer: M13 Forward
 POLYA=Yes.
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 Location/Qualifiers
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 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:2733092"
 /lab_host="DH10B (Life Technologies)"
 /clone_lib="NCI_CGAP_Sub6"
 /note="Vector: pT73E-Pac (Pharmacia) with a modified
 polylinker; Site 1: Not I; Site 2: Eco RI; NCI CGAP Sub6
 is a subtracted library derived from BM which consists of
 a mixture of four normalized libraries: NCI_CGAP_Brn5C,
 NCI_CGAP_Brn3, NCI_CGAP_Ov18, GSC1. The NCI_CGAP_Sub6
 library had 7 million recombinants. A single-stranded DNA
 preparation of BM was used as a tracer in a subtractive
 hybridization with a driver comprising the IMAGE pool
 (NCI CGAP Kid3 pool 1, LLAV 3334-3337, 3682-3683,
 3798-3803, IMAGE Clones 1323176-1323911,
 1456038-1456775, 1500552-1502855); NCI CGAP Kid5 pool 1:
 LLAV 3338-3342, 3722-3725, 3776-3778, IMAGE Clones 133912-1325831, 1471969-1472933, 1432-04-14932531;
 NCI CGAP L45 pool 1 LLAM 3575-3582, 385-3854, IMAGE
 Clones 1414920-1417991, 1520904-15224391, NCI CGAP GC4
 pool 1, LLAM 3164-3167, 3716-3720, 3733-3735, IMAGE
 Clones 1257096-1258631, 1463064-1470983, 1475592-1476743
 1; NCI CGAP Br22 pool 1 LLAM 2457-2459, 2758-2759,
 3042-3068, IMAGE Clones 985608-986759, 1101192-1101959,
 1217988-12206151; NCI CGAP GC1C pool 1 LLAM 2644-2653,
 2871-2872, IMAGE Clones 1057416-1061255, 114584-1145351;
 1; (50% of the driver population), plus a pool of 3,840
 arrayed clones from NCI_CGAP_Sub6, IMAGE Clones
 2708646-2710535 and NCI_CGAP_Sub2 (IMAGE Clones
 2710536-2712455) (20% of the driver population), plus a
 pool of 11,136 clones from NCI CGAP Sub3 (IMAGE Clones
 2712456-2723591) (30% of the driver population).
 Subtraction was performed as previously described [Bonaldi
 , Lennon & Soares (1996) : Normalization and Subtraction:
 Two Approaches To Facilitate Gene Discovery. Genome
 Research 6, 791-806].
 TAG LIB=NCI CGAP_Brn50
 TAG TISSUE=Brain
 TAG_SEQ=TTTCG"

```

BASE COUNT      194 a      53 c      77 g      172 t
ORIGIN
Query Match      51.1%; Score 511.2; DB 9; Length 532;
Best Local Similarity 99.4%; Pred. No. 1.9e-06;
Matches 513; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 391 TATATTAGTATAGCATTATTAAGGTTCTTGTGTGAGAGATCAGATAGAAATATATT 450
DB 17 TAAATAGTATAGCATTATTAAGGTTCTTGTGTGAGAGATCAGATAGAAATATATT 76
QY 451 TAAATGGTGGACATATTTCTAAGATACATACAGGATATTTTATACATAGAA 510
DB 77 TAAATGGTGGACATATTTCTAAGATACATACAGGATATTTTATACATAGAA 136
QY 511 ACAGAGAGTATTTACTTATTCATATTCATATTCGTTTACACATATACCTTTGATCAT 570
DB 137 ACAGAGAGTATTTACTTATTCATATTCATATTCGTTTACACATATACCTTTGATCAT 196
QY 571 TGATTTGTCATGCTTTTAAATATTCAGATAGAAATATTCGATTTACT 630
DB 197 TGTATGTCATGCTTTTAAATATTCAGATAGAAATATTCGATTTACT 256
QY 631 GAATCCTTGAAATAAATAAGCGGTCAGGTCAGCCTCAAACTTTAGAAAGCGTATACAGT 690
DB 257 GAATCCTTGAAATAAATAAGCGGTCAGGTCAGCCTCAAACTTTAGAAAGCGTATACAGT 716
QY 691 TGGACACAGGACTTTGGAGATACCACTGTTTAAAGGAATACCTTTGTAAGCTTACAGAAC 750
DB 377 TGGACACAGGACTTTGGAGATACCACTGTTTAAAGGAATACCTTTGTAAGCTTACAGAAC 776
QY 751 ATTATTACTTAAAGAGAGAAACACAGATGTTTCAATGACGTCATGCGGTAGAGAAAGGC 810
DB 377 ATTATTACTTAAAGAGAGAAACACAGATGTTTCAATGACGTCATGCGGTAGAGAAAGGC 836
QY 811 CTAACTTTACAGGACTTTTAAAGGCTTTACATATTTGGACCAAACTTGAAGATGAACC 870
DB 437 CTAACTTTACAGGACTTTTAAAGGCTTTACATATTTGGACCAAACTTGAAGATGAACC 936
QY 871 AGAAACAGAGACTTAAACAAATATACATGCAATG 906
DB 497 AGAAACAGAGACTTAAACAAATATACATGCAATG 932

RESULT 3
BU853405
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
EST.
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 743)
NIH-MGC http://mgi.nci.nih.gov/
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgabs@mail.nih.gov
Tissue Procurement: CLONTECH
CDNA Library Preparation: CLONTECH Laboratories, Inc.
DNA Sequencing by: Agencourt Bioscience Corporation
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LCC2870 row: a column: 13
High quality sequence stop: 648.
Location/Qualifiers
a . 743

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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6620245"
/lab_host="NIH MGC 82"
/clone_lib="NIH MGC 82"
/notes="Organ: testis; Vector: pDNR-LIB (Clontech); Site: 1;
Site 1: 199ccgctggcc; Site 2: 5' (ggccattaggcc); 5' and
3' adaptors were used in cloning as follows: 5' adaptor
sequence: 5'-CAGCGCATTTAGCC-3' and 3' adaptor sequence:
5'-ATTCTAGAGCCGAGCGCCGACATG-dT(30)BN-3'. Where B = A,
C, or G and N = A, C, G, or T. Average insert size
1.35 kb (range 0.9-4.0 kb). 14/15 colonies contained
inserts by PCR. This library was enriched for full-length
clones and was constructed by Clontech Laboratories (Palo
Alto, CA)."
BASE COUNT      248 a      123 c      121 g      250 t
ORIGIN
Query Match      49.3%; Score 403; DB 13; Length 743;
Best Local Similarity 100.0%; Pred. No. 0.0012;
Matches 403; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 599 AGATGAGAAATATTCGATATTCGATTCGATTCGATTCGATTCGATTCGATTCGATTCG 658
DB 11 AGATGAGAAATATTCGATATTCGATTCGATTCGATTCGATTCGATTCGATTCGATTCG 70
QY 659 TCAGCCTCAAACTTTAGAGAGGTCATACAGTTCGAGCAAGGACTTTGAGATACAGCTG 716
DB 71 TCAGCCTCAAACTTTAGAGAGGTCATACAGTTCGAGCAAGGACTTTGAGATACAGCTG 730
QY 719 TTTAAGGATATCTTTGTAAGCTTCGAGCAATTTTCTTAAAGAGGAAACAGAGAT 776
DB 121 TTTAAGGATATCTTTGTAAGCTTCGAGCAATTTTCTTAAAGAGGAAACAGAGAT 190
QY 779 CTCAATGAACGTCATCGGCTACAGAAACAGCTTAAGTTTACAGGACTTTTAGAGCTCT 832
DB 191 CTCAATGAACGTCATCGGCTACAGAAACAGCTTAAGTTTACAGGACTTTTAGAGCTCT 250
QY 839 ACATATTGTGCACCAAACTTGAAGATGAACAGAGAAACAGACTTAAACAAATATACAA 898
DB 251 ACATATTGTGCACCAAACTTGAAGATGAACAGAGAAACAGACTTAAACAAATATACAA 910
QY 899 TGCAAATGTAATTTTGTGTTTAAAGGCTTCGCTGATGTCACAGTATACCCAAATGG 958
DB 311 TGCAAATGTAATTTTGTGTTTAAAGGCTTCGCTGATGTCACAGTATACCCAAATGG 976
QY 959 ACATTAAGTTAGAGCAACAAACAAACCTGATTCGTCTTCTTT 1001
DB 371 ACATTAAGTTAGAGCAACAAACAAACCTGATTCGTCTTCTTT 413

RESULT 4
BU193776
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
EST.
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 877)
NIH-MGC http://mgi.nci.nih.gov/
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgabs@mail.nih.gov
Tissue Procurement: ATCC/DCTD/DTF
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

```

DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LINC at:
<http://image.llnl.gov>
Plate: LLAM3512 row: p column: 07
High quality sequence stop: 579.
Location/Qualifiers
1..877
/organism="Homo sapiens"
/mol_type="cDNA"
/db_xref="taxon:9606"
/clone="IMAGE:616142"
/tissue_type="melanocytic melanoma"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC 72"
/notes="Organ: skin; Vector: pCMV-SF016; Site 1: No.1;
Site 2: Salt; Cloned unidirectionally. Primer: Cligo dr.
Average insert size 2 kb. Library constructed by Life
Technologies."

BASE COUNT 300 a 166 c 177 g 234 t

ORIGIN
Query Match 39.8%; Score 396.2; DB 13; Length 877;
Best Local Similarity 99.3%; Pred. No. 0.0015; 3; Indels 0; Gaps 0;
Matches 400; Conservative 0; Mismatches 3;
QY 599 AGATGAGAAATATTCGATTATCTGACCTCAGTCAATCTTGAAATAAATAAAACGCGG 659
DB 293 AGATGAGAAATATTCGATTATCTGACCTCAGTCAATCTTGAAATAAATAAAACGCGG 352
QY 659 TCAGCCCTCAAACTTTAGAACGCTACATAGCTTGGACAGAGACTTTGAGATACCATG 718
DB 353 TCAGCCCTCAAACTTTAGAACGCTACATAGCTTGGACAGAGACTTTGAGATACCATG 412
QY 719 TTATAGGAAATACCTTTGTAACCTGACAGAACTTTACTTAAAGAGGAAACACAGAT 778
DB 413 TTATAGGAAATACCTTTGTAACCTGACAGAACTTTACTTAAAGAGGAAACACAGAT 472
QY 779 CTTCAATGAAAGCTATCGGCTACAGAAAGCGCTTAAGTTTACAGGACTTTTACAGTCT 838
DB 473 CTTCAATGAAAGCTATCGGCTACAGAAAGCGCTTAAGTTTACAGGACTTTTACAGTCT 532
QY 839 ACATATTGCGCAAACTTGAAGTGAAGCGAGAAACAGACTTAAACAATAACAA 898
DB 533 ACATATTGCGCAAACTTGAAGTGAAGCGAGAAACAGACTTAAACAATAACAA 592
QY 899 TGCAGATGTAATTTTGTGTTTAAAGGCTTGCCTTGTGTCACAGTATCCCAATG 958
DB 593 TGCAGATGTAATTTTGTGTTTAAAGGCTTGCCTTGTGTCACAGTATCCCAATG 652
QY 959 ACCTAAGTTAGAGCACAAACAAACCTGATCTGGCTCTTTT 1001
DB 653 ACCTAAGTTAGAGCACAAACAAACCTGATCTGGCTCTTTT 695

RESULT 5
EG204678
LOCUS EG204678 784 bp mRNA linear EST 21-APR-2001
DEFINITION R5724093 Athersys RAGE Library Homo sapiens cDNA, mRNA sequence.
ACCESSION EG204678
VERSION EG204678.1 GI:13726365
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 784)
REFERENCE
AUTHORS Harrington, J.C., Sherf, B., Rundlett, S., Jackson, P.D., Perry, R.,
Cain, S., Leventhal, C., Thornton, M., Ramachandran, R., Harrington, J.,
Lerner, L., Costanzo, D., Xceligott, K., Booser, S., Mays, R., Smith,
E., Veloso, N., Klika, A., Hess, J., Cothren, K., Lo, K., Offenbacher,
J., Danzig, J. and Ducar, M.
Creation of genome-wide protein expression libraries using random.

activation of gene expression
Nat. Biotechnol. 19 (5): 440-445 (2001)
21227151
PUBMED
11329013
COMMENT
Contact: Scott J. Cain
Athersys, Inc. 4401
3201 Carnegie Ave, Cleveland, OH 44115, USA
Tel: 216 431 9900
Fax: 216 361 9596
Email: scain@atersys.com
High quality sequence stop: 500.
Location/Qualifiers
1..784
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/cell_line="HT1080"
/clone_lib="Athersys RAGE Library"
/notes="See 'Creation of Genome-wide Protein Expression
Libraries using Random Activation of Gene Expression',
Nature Biotechnology, in press. Note that even though the
cell type indicated is HT1080, since a random activation
method was used, these sequence tags are not necessarily
expressed in HT1080 under normal circumstances."
BASE COUNT 255 a 141 c 129 g 259 t
ORIGIN
Query Match 38.8%; Score 388.6; DB 10; Length 784;
Best Local Similarity 99.0%; Pred. No. 0.0029; 4; Indels 0; Gaps 0;
Matches 391; Conservative 0; Mismatches 4;
QY 607 AATATTGATTATCTGACCTCAGTCAATCTTGAAATAAATAAAACGCGCTCAGCCCT 666
DB 7 AATATTGATTATCTGACCTCAGTCAATCTTGAAATAAATAAAACGCGCTCAGCCCT 66
QY 667 CAATCTTTAGAACGCTACATAGCTTGGACAGAGACTTTGGAGATACCACTTTTAAAGGA 726
DB 67 CATCTTTAGAACGCTACATAGCTTGGACAGAGACTTTGGAGATACCACTTTTAAAGGA 126
QY 727 AATACCTTTGTAACCTTGCAGAACACTTTACTTAAAGAGAGAAACAGAAATCTTCAATG 766
DB 127 AATACCTTTGTAACCTTGCAGAACACTTTACTTAAAGAGAGAAACAGAAATCTTCAATG 186
QY 767 AACGCTATCGCTACAGAAACAGGCTTAGTTTACAGGACTTTTAGAGCTCTTACATATTT 846
DB 187 AACGCTATCGCTACAGAAACAGGCTTAGTTTACAGGACTTTTAGAGCTCTTACATATTT 246
QY 847 GTGACCAAACTTGAAGATGAACAGAGAAACAGACTTAAACAAATATACAAATGCAATG 906
DB 247 GTGACCAAACTTGAAGATGAACAGAGAAACAGACTTATACAAATATACAAATGCAATG 306
QY 907 TAAATTTTGTGTTTAAAGGCTTGCCTTGTGTCACAGTATCCCAATGCACTAAG 966
DB 307 TAAATTTTGTGTTTAAAGGCTTGCCTTGTGTCACAGTATCCCAATGCACTAAG 366
QY 967 TTAGAGCACAAACAAACCTGATCTGGCTCTTTT 1001
DB 367 TTAGAGCACAAACAAACCTGATCTGGCTCTTTT 401

RESULT 6
EG217901
LOCUS EG217901 749 bp mRNA linear EST 21-APR-2001
DEFINITION R5737625 Athersys RAGE Library Homo sapiens cDNA, mRNA sequence.
ACCESSION EG217901
VERSION EG217901.1 GI:13743922
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 749)
REFERENCE
AUTHORS Harrington, J.C., Sherf, B., Rundlett, S., Jackson, P.D., Perry, R.,

Cain, S., Leventhal, C., Thornton, M., Ramachandran, R., Whittington, J., Lerner, J., Costanzo, D., McElligott, K., Booser, S., Mays, R., Smith, E., Veloso, N., Klika, A., Hess, J., Cochren, K., J.C.K., Offenbacher, C., Danzig, J., and Ducar, X.

TITLE
Creation of genome-wide protein expression libraries using random activation of gene expression

JOURNAL
Nat. Biotechnol. 19 (5), 440-445 (2001)

MEDLINE
21227151

PIBMED
11329013

COMMENT
Contact: Scott J. Cain
Athersys, Inc.
3201 Carnegie Ave., Cleveland, OH 44115, USA
Tel: 216 431 9900
Fax: 216 361 9596
Email: scain@atersys.com
High quality sequence stop: 551.

FEATURES
source
1..749
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/cell_line="H1080"
/clone_lib="Athersys RAGE Library"
/note="See 'Creation of Genome-wide Protein Expression Libraries using Random Activation of Gene Expression', Nature Biotechnology, in press. Note that even though the cell type indicated is H1080, since a random activation method was used, these sequence tags are not necessarily expressed in H1080 under normal circumstances."

BASE COUNT
251 a 127 c 121 g 250 t

ORIGIN
Query Match 38.8%; Score 389; DB 10; Length 743;
Best Local Similarity 99.1%; Pred. No. 3.0031;
Matches 399; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

603 GAGAAATATTCGATTATC TGATTTCACTGGAATCTTGAAAAAATAAAGCAGCTCA 661
|||||
1 GAGAAATATTCGATTATC TGATTTCACTGGAATCTTGAAAAAATAAAGCAGCTCA 60
|||||

662 GCGCTCAAACTTTAGAGCGTACATCGTTTGAACAAGGACTTTGGAGATACCAGTCTT 721
|||||

61 GCGCTCAAACTTTAGAGCGTACATCGTTTGAACAAGGACTTTGGAGATACCAGTCTT 120
|||||

722 AAGGAATACCTTTGTAACCTTGGAGACATTTTACTTAAAGAGGAACACACAGATCTT 781
|||||

121 AAGGAATACCTTTGTAACCTTGGAGACATTTTACTTAAAGAGGAACACACAGATCTT 180
|||||

782 CAATGAACGCTCATCGGCTACAGAAACAGGCTTAAGTTTACAGGACTTTTAGAGCTTTACA 841
|||||

181 CAATGAACGCTCATCGGCTACAGAAACAGGCTTAAGTTTACAGGACTTTTAGAGCTTTACA 240
|||||

842 TATTTGTGCACCAACTTGAAGATGAACCCAGAAACAGACTTTAAACAAATATACAAATGC 901
|||||

241 TATTTGTGCACCAACTTGAAGATGAACCCAGAAACAGACTTTAAACAAATATACAAATGC 300
|||||

902 AATGTAAATTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTT 961
|||||

301 AATGTAAATTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTT 360
|||||

962 CTAAGTTAGGACACAAACAACTGATTCGGTCTCTCTT 1001

361 CTAAGTTAGGACACAAACAACTGATTCGGTCTCTCTT 400
|||||

RESULT 7
BG193969/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
BG193969
BG193969
EST
Homo sapiens (human)

894 bp mRNA linear EST 21-APR-2003
Athersys RAGE Library Homo sapiens cDNA, mRNA sequence.

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
1 (bases 1 to 894)
Harrington, J.J., Sherf, B., Rundlett, S., Jackson, P.D., Perry, R., Cain, S., Leventhal, C., Thornton, M., Ramachandran, R., Whittington, J., Lerner, J., Costanzo, D., McElligott, K., Booser, S., Mays, R., Smith, E., Veloso, N., Klika, A., Hess, J., Cochren, K., J.C.K., Offenbacher, C., Danzig, J., and Ducar, X.

TITLE
Creation of genome-wide protein expression libraries using random activation of gene expression

JOURNAL
Nat. Biotechnol. 19 (5), 440-445 (2001)

MEDLINE
21227151

PIBMED
11329013

COMMENT
Contact: Scott J. Cain
Athersys, Inc.
3201 Carnegie Ave., Cleveland, OH 44115, USA
Tel: 216 431 9900
Fax: 216 361 9596
Email: scain@atersys.com
High quality sequence stop: 461.

FEATURES
source
1..894
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/cell_line="H1080"
/clone_lib="Athersys RAGE Library"
/note="See 'Creation of Genome-wide Protein Expression Libraries using Random Activation of Gene Expression', Nature Biotechnology, in press. Note that even though the cell type indicated is H1080, since a random activation method was used, these sequence tags are not necessarily expressed in H1080 under normal circumstances."

BASE COUNT
284 a 154 c 169 g 283 t 4 others

ORIGIN
Query Match 35.0%; Score 350; DB 10; Length 894;
Best Local Similarity 96.1%; Pred. No. 0.029;
Matches 389; Conservative 0; Mismatches 12; Indels 3; Gaps 3;

599 AGATGAGAAATATTCGATTATTCGACTTCACGTGATCTTGAAAAAATAAAGCAGCG 658
|||||

770 AGATGAGAAATATTCGATTATTCGACTTCACGTGATCTTGAAAAAATAAAGCAGCG 711
|||||

659 TCAGCCCTCAAACTTT-AGAAGCGTACATACGTGGAAACAGGACTTTGGAGATACCAC 717
|||||

710 TCAGCCCTCAAACTTT-AGAAGCGGACATACGTGGAAACAGGACTTTGGAGATACCAC 651
|||||

718 GTTTAAGGAATACCTTTGTAACCTTCCAGACATTTACTTAAAGAGGAACACACAGA 777
|||||

650 GTTTAAGGAATACCTTTGTAACCTTCCAGACATTT-CTTAAAGAGGAACACACAGA 593
|||||

778 TCITCAATGAACGCTCATCGGCTACAGAAACAGGCTTAAGTTTACAGGACTTTTAGAGTCT 837
|||||

592 TCITCAATGAACGCTCATCGGCTACAGAAACAGGCTTAAGTTTACAGGACTTTTAGAGTCT 533
|||||

838 TACATATTTGTGCACCAAACTTGAAGATGAACCCAGAAACAGACTTTAAACAAATATACA 897
|||||

532 TACATATTTGTGCACCAAACTTGAAGATGAACCCAGAAACAGACTTTAAACAAATATACA 473
|||||

896 ATGCAATGTAAATTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTT 957
|||||

472 ATGCAATGTAAATTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTT 413
|||||

956 GACACTAAGTTAGGACACAAACAACTGATTCGGTCTCTCTT 1001
|||||

412 GACACTAAGTTAGGACACAAACAACTGATTCGGTCTCTCTT 369
|||||

RESULT 8
BF216343
LOCUS
BF216343
728 bp mRNA linear EST 26-NOV-2000

DEFINITION 601884376F1 NIH_MGC_57 Homo sapiens cDNA clone IMAGE:410722 5',
mRNA sequence.
ACCESSION BF216343
VERSION BF216343.1 GI:11109329
KEYWORDS EST
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 728)
NIH-MGC <http://mimc.nci.nih.gov/>
National Institutes of Health, Mammalian Gene Collection (MGC).
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs@nci.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: CLONTECH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LCM976 row: h column: 19
High quality sequence stop: 574
Location/Qualifiers
1..728
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:410722"
/issue_type="glioblastoma"
/lab_host="DH10B (1:1) (page-resistant)"
/clone_1b="NIH_MGC_57"
/notes="Organ: brain; Vector: pDRF-lib (Clontech). Site: 1:
5'UTR (ggcgctcgcc); Site 2: 5'UTR (ggcgctcgcc);
Double-stranded cDNA was prepared from cell line RNA. 5'
and 3' adaptors were used in cloning as follows. 5'
adaptor sequence: 5'-CACGGCATTTGGCC-3' and 3' adaptor
sequence: 5'-ATTCTAGAGCGGAGCGCGGACAG-G-ATTCTTBN-3'
(where B = A, C, G and N = A, C, G, or T). Average
insert size 1.55 kb (range 0.9-4.0 kb). 12715 colonies
contained inserts by PCR. This library was enriched for
full-length clones and was constructed by Clontech
Laboratories (Palo Alto, CA).
BASE COUNT 248 a 132 c 150 g 198 t
ORIGIN
Query Match 28.6%; Score 286.12; DS 10; Length 728;
Best Local Similarity 92.6%; Pred. No. 1.8;
Matches 377; Conservative 0; Mismatches 23; Indels 7; Gaps 7;
CY 599 AGATGAGAAATATCGATTATCTGACTTACTGAACTCTGMAAAATAAAGCGAGCG 456
Db 142 AGATGAGAAATATCGATTATCTGACTTACTGAACTCTGMAAAATAAAGCGAG-G 269
CY 659 TCAGCGCTCAAACTTACAGCGCTACATCGTTGGACACAGGACTT-TGGAGATACACT 317
Db 201 TCAGCGCTCAAA-TTTAGACGCTACATCGTTGGACACAGGACTT-GTGAGATACACT 259
CY 718 -GTTTAGGAAATACCTT-TGTAACTCGAGAACATTTTACTTAAAGAGGAAACACAA 775
Db 260 GGTTTAGGAAATACCTTGTGTAACTCGAGAACATTTTACTTAAAGAGGAAACACAA 319
CY 776 GATCTTCAATGACGCTATCGGCTACAGAAACAGCGCTAAGTTTACAGGATTTTATAGT 835
Db 320 GATCTTCAATGACGCTATCGGCTACAGAAACAGCGCTAAGTTTACAGGATTTTATAGT 379
CY 836 CTTTACATTTTGCACCAACTTGAGATGACCGAGAAACACAGCTTAAACAAATATA 895
Db 380 CTTTACATTTTGCACCAAACTTGAGATGACCGAGAAACACAG-TTAAACAAATATA 438
CY 896 CAATGCAATGTAATTTTGTGTTTAAAGSCCTTGCCTTGTGTTGTCACAGTTATCCCA 955

Db 439 CAATGCAATGTAATTTTGTGTTTAAAGSCCTTGCCTTGTGTTGTCACAGTTATCCCA 498
CY 356 TGGACACTAAG-TTAGAGCACACAAAACTGATTTGTGTTGTTT 1001
Db 499 TGGACACTAAGTTTAGAGCACACAAAACTGATTTGTGTTGTTT 545
RESULT 3
BG185583 360 bp mRNA linear EST 21-APR-2001
DEFINITION R574534 Athysys RAGE Library Homo sapiens cDNA, mRNA sequence.
ACCESSION BG185583
VERSION BG185583.1 GI:13707270
KEYWORDS EST
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 360)
Harrington, J.J., Sherf, B., Rundle, S., Jackson, P.D., Perry, R.,
Cain, S., Leventhal, C., Thornton, M., Raghavendran, R., Whittington, J.,
Lerner, L., Costanzo, D., McEligott, K., Booser, S., Mays, R., Smith,
E., Veloso, N., Klika, A., Hess, J., Cothren, K., Lo, K., Offenbacher,
J., Danzig, J. and Ducar, M.
Creation of genome-wide protein expression libraries using random
activation of gene expression
Nat. Biotechnol. 19 (5), 440-445 (2001).
JOURNAL Nat. Biotechnol.
MEDLINE 21227151
PubMed 11329013
COMMENT Contact: Scott J. Cain
Athersys, Inc.
320 Carnegie Ave., Cleveland, OH 44115, USA
Tel: 216 431 9900
Fax: 216 361 9596
Email: scain@atersys.com
High quality sequence stop: 310.
Location/Qualifiers
1..360
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_1b="HT1080"
/clone_2b="Athersys RAGE Library"
Libraries using Random Activation of Gene Expression,
Nature Biotechnology, in press. Note that even though the
cell type indicated is HT1080, since a random activation
method was used, these sequence tags are not necessarily
expressed in HT1080 under normal circumstances.
BASE COUNT 129 a 61 c 53 g 107 t
ORIGIN
Query Match 27.5%; Score 277.8; DS 10; Length 360;
Best Local Similarity 97.1%; Pred. No. 5;
Matches 304; Conservative 0; Mismatches 7; Indels 2; Gaps 2;
CY 659 GTTGAAACAGGACTTTGGAGATACCACTGTTAAGGAATACCTTTGTAAAGCTGCGA 748
Db 15 GTTGAAACAGGACTTTGGAGAT-CCACTGTTGAAGGAATACCACTTTGAACCTGCGA 73
CY 749 ACATTTTACTTAAAGAGGAAACACAGATCTTCAATGAACGTCATCGGTACAGAAACA 808
Db 74 ACATTTTACTTAAAGAGGAAACACAGATCTTCAATGAACGTCATCGGTACAGAAACA 133
CY 809 GCTTAAGTTTACAGGACTTTTAGAGCTTACATATTGTGCACCAAACTTTGAGATGAA 569
Db 134 GCTT-AGTTTACAGGACTTTTAGAGCTTACATATTGTGCACCAAACTTTGAGATGAA 192
CY 869 CCAGAAACACAGCTTAAACAAATATACAAATGCAATGTAATTTTGTGTTTAAAGGCC 928
Db 193 CCAGAAACACAGCTTAAACAAATATACAGTGCAGTAATTTTGTGTTTAAAGGCC 252
CY 929 TTGCGCTTATGCTCACAGTTATCCCAATGGACACTTAAGTTAGAGCACCAAAACCTCAT 988

```

Db 253 TGGCTTATGTCACACTTATCCCAATGGACACTAGTATAGACACACACAAACCTGAT 312
QY 989 TCTGCTCTCTTT 1001
Db 313 TCTGCTCTCTTT 325

RESULT 10
BE172740 460 bp mRNA linear EST 21-APR-2001
LOCUS XRO-H70559-123400-010-309 H70559 Homo sapiens cDNA, mRNA sequence.
DEFINITION BE172740
ACCESSION BE172740
VERSION BE172740.1 GI:8635466
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
REFERENCE 1 (bases 1 to 460)
AUTHORS Dias Neto,E., Garcia Correa,R., Vertovski-Rimeida,S., Stienos,M.R.,
Nagai,M.A., da Silva,W.C., Zago,M.A., Bordin,S., Costa,F.P.,
Goldman,G.H., Carvalho,A.F., Matsukura,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., de Oliveira,P.S., Bucher,P., Congenel,C.V., O'Hare,
M.J., Soares,F., Brennan,R.R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.
TITLE Shotgun sequencing of the human transcriptome with 2RF expressed
sequence tags
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
MEDLINE 20202663
PUBMED 10737800
COMMENT Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01500-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the RAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=&t2=MR0-H70559-120
400-010-a09&t3=2000-04-12&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 6
High quality sequence stop: 460.
Location/Qualifiers
1..460
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="H70559"
/notes="Organ: head/neck; Vector: puc18; Site: 1. Small;
Site 2: Small; A mini-library was made by cloning products
derived from CRESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."
BASE COUNT 117 a 112 c 94 g 147 t
ORIGIN
Query Match 24.4% Score 244.6; DB 10; Length 460;
Best Local Similarity 74.6% Pred. No. 33;
Matches 326; Conservative 0; Mismatches 99; Indels 12; Gaps 1;

QY 5 TAAAAATGCATCTTAGCTGGGCAAGTGGCTCACCTGTGTATCCACACTTTGGG 64
Db 460 TATATATATATAAAAGCCAGGTGGCGTGTTCACACCTGTATATCCACACTTTGGG 401
QY 65 AGGCCAAGTGGCGCATCTACCTAGGTAGAGTTCCAGGACCAACCTGGCATCTGGC 124

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Db 400 AGCCACAGACAGAGGTGGATCATAGGTCAGAGTTCAGAGACCGCTGGCCACAGTGGT 341
QY 125 GAAACCTGTCTCTACTATAAATAACAAAATTAGCTGGGCATGGAGGCAATGTCCTGTAA 184
Db 340 GAAACCTGCCTCTACTATAAATAACAAAATTAGCTAGGCGTGGTGGGAGGAGCTGTAA 281
QY 185 TCCAGCTACTCGGAGAGCTGAGACAGAGATTCTTGMACCTTGGAGGGGGAGGTGGC 244
Db 280 TCCAGCTACTAGGAGGTGGAGGAGGAGATCACTTGAATCTGGAGGAGAGATGGC 221
QY 245 ATATCTGAGTGTGAAATGTGATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 292
Db 210 ANTGAGCCGAGGTGCACACTTGCACCTCCAGCTGGGCGACAGAGATGAAATCTGCTC 161
QY 293 TTGAACTTTTATATAATGATGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 352
Db 160 AAAAAATATATATAAATGATGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 101
QY 353 AATGTGTATCTATGAACATAACACAGTAATAATATATATATATATATATATATATAT 412
Db 100 AATGTGTATCTATGAACATAACACAGTAATAATATATATATATATATATATATAT 41
QY 413 AGGTTCTTGTGTAGCA 429
Db 40 AGGTTCTTGTGTAGCA 24

RESULT 11
LOCUS RG208384
DEFINITION RST27882 Athersys RAGE Library Homo sapiens cDNA, mRNA sequence.
ACCESSION RG208384
VERSION RG208384.1 GI:13730071
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
REFERENCE 1 (bases 1 to 239)
AUTHORS Harrington,J.J., Sherif,B., Rundlett,S., Jackson,P.D., Perry,R.,
Cain,S., Jevencal,C., Thornton,M., Ramachandrar,R., Whittington,S.,
Lerner,L., Costanzo,D., McEligott,K., Booser,S., Mays,R., Smith,
E., Veloso,N., Kiika,A., Hess,J., Cothren,K., Lo,K., Offenbacher,
J., Danzig,J. and Ducar,M.
TITLE Creation of genome-wide protein expression libraries using random
activation of gene expression
JOURNAL Nat. Biotechnol. 19 (5), 440-445 (2001)
MEDLINE 21227151
PUBMED 11329013
COMMENT Contact: Scott J. Cain
Athersys, Inc
3201 Carnegie Ave. Cleveland, OH 44115, USA
Tel: 216 431 9900
Fax: 216 361 9596
E-mail: scaina@atersys.com
High quality sequence stop: 189.
Location/Qualifiers
1..239
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_lib="Athersys RAGE Library"
/notes="See 'Creation of Genome-wide Protein Expression
Libraries using Random Activation of Gene Expression',
Nature Biotechnology, in press. Note that even though the
cell type indicated is HT1080, since a random activation
method was used, these sequence tags are not necessarily
expressed in HT1080 under normal circumstances."
BASE COUNT 89 a 46 c 42 g 63 t
ORIGIN
Query Match 21.8% Score 218.2; DB 10; Length 239;

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Best Local Similarity 96.5%; Pred. No. 2,8e+02;
Yatches 723; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

QY 656 GCGTGGAGCCCTCAAACTTTAGAGCGGTACATACGTTGGAACAGAGACCTTTGAGATACCA 715
DB 5 GTGTAGCCCTCATTTTTTAGAGCGGTACATACGTTGGAACAGAGACCTTTGAGATACCA 67

QY 716 CTGTTTAAAGAAATACCTTTGTAACCTTGCAGAACATTTTCTTAAAGAGAGAACACAA 775
DB 6 CTGTTTAAAGAAATACCTTTGTAACCTTGCAGAACATTTTCTTAAAGAGAGAACACAA 127

QY 776 GATCTTCATGATGATCATGCGGTACAGAAACAGGCTAGTTTACAGACATTTTACAGT 835
DB 128 GATCTTCATGATGATCATGCGGTACAGAAACAGGCTAGTTTACAGACATTTTACAGT 167

QY 836 CTTCATATTTGTGCACCAACTTGAAGATGAACAGAAACAGACTTAA 886
DB 188 CTAACTATTTGTGCACCAACTTGAAGATGAACAGAAACAGACTTAA 236

RESULT 12
LOCUS W42916 401 bp mRNA linear EST 10-OCT-1996
DEFINITION ZC25f07.x1 Soares senescent fibroblasts NbHSP Homo sapiens cDNA
Clone IMAGE:323365, similar to SW:YEN7_YEAST P39955 HYPOPHOSPHATASE
100-3 KC PROTEIN IN ME14-CAU1 INTERGENIC REGION. (1); mRNA
sequence.
ACCESSION W42916 GI:1327407
VERSION W42916
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 401)
AUTHORS Buller, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman,
M., Huftman, W., Kucaba, T., Le, M., Lennon, G., Manta, W., Parsons, J.,
Ridkin, J., Rohlfing, T., Soares, M., Tan, P., Trivaskis, E., Waterson,
J.R., Williamson, A., Wolfdmann, P. and Wilson, R.
TITLE The WashU-Merck EST Project
JOURNAL Unpublished
COMMENT Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63128
Tel: 314 286 1500
Fax: 314 286 1810
Email: estswatson.wustl.edu
This clone is available royalty-free through LNCX; contact the
IMAGE Consortium (info@image.lln.gov) for further information.
Insert Length: 2552 Std Error: 0.00
Seq primer: mob.R3GA-E7
High quality sequence stop: 291.
FEATURES
Location/Qualifiers
1..401
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="Gene:1254577"
/db_xref="taxon:9606"
/clone="IMAGE:323365"
/tissue_type="senescent fibroblast"
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares senescent fibroblasts NbHSP"
/notes="Vector: pT73D (Pharmacia) with a modified
polylinker V-type; phagemid; Site 1: Not 1; Site 2: Eco R1;
1st strand cDNA was primed with a Not 1 - oligo(dT)
primer (5'
TGTACCAATCTGAGTGGAGCGCGGCATTTTTTTTTTTTTTTT 3';
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pT73 vector
(Pharmacia). Library went through one round of
normalization to a Cot = 5. Library constructed by Bento
Soares and X. Fatima Bonaldo."
BASE COUNT 87 a 145 c 118 g 166 t
ORIGIN
Query Match 19.0%; Score 190.4; DB 28; Length 516;
Best Local Similarity 87.1%; Pred. No. 8.8e-02;
Yatches 209; Conservative 0; Mismatches 31; Indels 3; Gaps 0;

BASE COUNT 140 a 75 c 79 g 95 t 10 others
ORIGIN
Query Match 20.9%; Score 209.4; DB 14; Length 401;
Best Local Similarity 96.3%; Pred. No. 3.3e+02;
Mismatches 20; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 999 AGATGAGAATATTTCGATTATCTGATCTTCACTGAATCTCTTGAAAAAATAAAGCGAGCG 958
DB 163 AGATGAGAATATTTCGATTATCTGATCTTCACTGAATCTCTTGAAAAAATAAAGCGAGCG 242

QY 659 TCAGCCCTCAAACTTTAGAGCGGTACATACGTTGGAACAGAGACCTTTGAGATACCACTG 718
DB 243 TCAGCCCTCAAACTTTAGAGCGGTACATACGTTGGAACAGAGACCTTTGAGATACCACTG 302

QY 719 TTTAAGGAAATACCTTTGTAACCTTGCAGAACATTTTCTTAAAGAGAGAACACAAAGAT 778
DB 303 TTTAAGGAAATACCTTTGTAACCTTGCAGAACATTTTCTTAAAGAGAGAACACAAAGAT 362

QY 779 CTTCATGATGATCATGCGGTACAGAAACAGGCTAGTTTACAGACATTTTACAGT 816
DB 363 CTTCATGATGATCATGCGGTACAGAAACAGGCTAGTTTACAGACATTTTACAGT 400

RESULT 13
LOCUS A012451 516 bp DNA linear GSS 29-AUG-1998
DEFINITION CIT-HSP-2372C9.TF CIT-HSP Homo sapiens genomic clone 2372C9,
genomic survey sequence.
ACCESSION A012451 GI:3484611
VERSION A012451
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 516)
AUTHORS Adams, X.D., Rounsley, S.D., Zhao, S., Bass, S., Linher, K., Golden, K.,
Berry, K., Granger, D., Suh, E., Wible, C., Shizuya, H., Simon, M. and
Venter, J.C.
TITLE Use of a random human BAC End Sequence Database for Sequence Ready
JOURNAL Unpublished
COMMENT Other_GSSs: CIT-HSP-2372C9.TF
Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 3200
Fax: 301 838 3308
Email: mcdams@tigr.org
Clones are available from Research Genetics (info@resgen.com). BAC
end search page:
http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html.
Seq primer: M13-2;
Class: BAC ends.
FEATURES
Location/Qualifiers
1..516
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="2372C9"
/sex="Male"
/cell_type="Sperm"
/notes="Vector: pBelOBAC11; Site 1: HindIII; Site 2:
HindIII"
BASE COUNT 87 a 145 c 118 g 166 t
ORIGIN
Query Match 19.0%; Score 190.4; DB 28; Length 516;
Best Local Similarity 87.1%; Pred. No. 8.8e-02;
Yatches 209; Conservative 0; Mismatches 31; Indels 3; Gaps 0;

```

```

QY 6 AAAATGCACTATTAGGCTGSSCAAACTGGCTCAGCTCTCTATATCCAGACACTTTGGA 65
DB 506 AAAAATGCAAAATTTAGCTGGGGCACTGGCCAGAGCTGTATATCCAGACACTTTGGA 447
QY 66 GGGCAAGTGGGGGATCAGCTGAGGTAGGAGTTCAGGACCACTGGGCAATGAGG 125
DB 446 GGGCAAGTGGGGGATCAGCTGAGGTAGGAGTTCAGGACCACTGGGCAATGAGG 387
QY 126 AAAACCTGCTCTACTAAAAATACAAAATTTAGTGGGCACTGGAGGAGTGTGCTGTAT 185
DB 386 AAAACCTGCTCTACTAAAAATACAAAATTTAGTGGGCACTGGAGGAGTGTGCTGTAT 327
QY 186 CCCGCTACTGGGAGCTGGAGAGGAGATTTCTTGAACCTGGAGGGGGAGGTGCA 245
DB 326 CCCGCTACTGGGAGCTGGAGAGGAGATTTCTTGAACCTGGAGGGGGAGGTGCA 267

CA946732 515 bp -RNA linear EST 31 DEC-2002
LOCUS CA946732
DEFINITION mRNA sequence.
ACCESSION CA946732
VERSION CA946732.1 GI:2743609
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 515)
Lenschka, J., Pearce, X., Brestelli, C., Gradwohl, G., Clifton, S.,
Hillier, L., Marto, M., Pape, D., Wille, F., Martin, J., Slight, A.,
Schmitt, A., Theising, P., Ratter, S., Rorko, I., Bennett, S., Cardenas,
N., Gibbons, K., McCann, R., Cole, R., Tsagaris, V., Williams, T.,
Jackson, Y., and Bowen, Y.
Endocrine Pancreas Consortium
Unpublished
Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
Endocrine Pancreas Consortium
Harvard University, Howard Hughes Medical Institute
Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
MA 02138
Tel: 617-495-1812
Fax: 617-495-8557
Email: dmelton@bioh.p.harvard.edu
Library was constructed by Dr. Hiroshi Inoue DNA sequencing by:
Washington University Genome Sequencing Center For information on
obtaining a clone please contact: Dr. Hiroshi Inoue
(hinoue@im.wustl.edu)
Seq primer: -40RP from Gibco
High quality sequence stop: 451.
Location/Qualifiers
1..515
/organism="Homo sapiens"
/mol_type="-RNA"
/db_xref="taxon:9606"
/clone="IMAGE:5164654"
/tissue_type="Purified pancreatic islet"
/lab_host="DH10B"
/clone_lib="HP85 islet"
/notes="Organ: Pancreas; Vector: pBlueScript SK(-); Site_1:
Not1; Site_2: Xho1; cDNA made by oligo-dT priming.
Size selected on agarose gel. Average insert size ~1kb. 5'
Xho1 site was destroyed after directional cloning.
Amplified once. Contact information: Hiroshi Inoue, MD,
Metabolism Div, (Alan Permut Lab), Washington University
School of Medicine, Box 8129, 660 South Euclid Ave., St.
Louis, MO 63110, E-mail: hinoue@im.wustl.edu, Tel:
314-362-1916, Fax: 314-747-2692."
94 a 154 c 114 g 153 t
BASE COUNT
ORIGIN

```

```

Query Match 12.9%; Score 188.8; DB 14; Length 515;
Best Local Similarity 85.7%; Pred. No. 9.8e-02;
Matches 238; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

QY 6 AAAATGCACTATTAGGCTGGGCAAACTGGCTCAGCTCTCTATATCCAGACACTTTGGA 65
DB 297 AAAATACAAAATTTAGGCTGGGCAAGTGGCCAGAGCTGTATATCCAGACACTTTGGA 238
QY 66 GGGCAAGTGGGGGATCAGCTGAGGTAGGAGTTCAGGACCACTGGGCAATGAGG 125
DB 237 GGGCAAGTGGGGGATCAGCTGAGGTAGGAGTTCAGGACCACTGGGCAATGAGG 176
QY 126 AAAACCTGCTCTACTAAAAATACAAAATTTAGTGGGCACTGGAGGAGTGTGCTGTAT 185
DB 177 AAAACCTGCTCTACTAAAAATACAAAATTTAGTGGGCACTGGAGGAGTGTGCTGTAT 118
QY 186 CCCGCTACTGGGAGCTGGAGAGGAGATTTCTTGAACCTGGAGGGGGAGGTGCA 245
DB 117 CCCGCTACTGGGAGCTGGAGAGGAGATTTCTTGAACCTGGAGGGGGAGGTGCA 58

RESULT 14
LOCUS AI1339725/c
DEFINITION 3942e06.x1 Soares NHPM4.S: Homo sapiens cDNA clone IMAGE:1935206
3 similar to contains Alu repetitive element; contains element PTF5
repetitive element 1; mRNA sequence.
ACCESSION AI1339725
VERSION AI1339725.1 GI:4076652
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 446)
NCI-CCGAP http://www.ncbi.nlm.nih.gov/ccgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: ccgaps@mail.nih.gov
This clone is available royalty-free through LNCX; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert length: 2084
Seq primer: -40UP from Gibco
High quality sequence stop: 446.
Location/Qualifiers
1..446
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1935206"
/tissue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"
/lab_host="DH10B"
/clone_lib="Soares NHPM4.S"
/notes="Organ: mixed (see below); Vector: pTZ193D-Eac
(parracina) with a modified polylinker; Site_1: Not 1;
Site_2: Eco RI; Equal amounts of plasmid DNA from three
normalized libraries (melanocyte 2NBHM, pregnant uterus
NBHPJ, and fetal heart NBHH19W) were mixed, and ss circles
were made in vitro. Following HAP purification, this DNA
was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from pools of
5,000 clones made from the same 3 libraries. The pools
consisted of 1 M.A.G.E. clones 260232-265223,
340488-345479, and 484488-489479."
120 a 117 c 99 g 110 t
BASE COUNT
ORIGIN

Query Match 18.7%; Score 187.4; DB 9; Length 446;
Best Local Similarity 85.3%; Pred. No. 1.2e-03;

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Matches	209;	Conservative	3;	Mismatches	36;	Indels	3;	Gaps	0;
QY	1	AAATATAAAATGCATACATTAGGCTGGCAAGTGGCTCACGCTGTSTANTCCACGACTT	60						
DB	300	AAATTATTAGAATTAAATCTGGCTGGGGGGCTGGGCAACGCTGTANTCCACGACTT	241						
QY	41	TGGGAGGCCAAGGTGGGGGATCACTGAGGTAGGATTAGGACCAAGCTGGGCACTCA	120						
DB	240	TGGGAAGCCCAAGGTGGGGGATCACTTCAAGTTAGGASTTGAAGACCAAGCTGGGCAACA	181						
QY	121	TGGCGAAGCCGTCTCTACTTAAATACAAAATTAGCTGGGCAATGAGGCACTGGGCT	180						
DB	180	TGGCGAAGCCGTCTCTACTTAAATACAAAATTAGCGGGGCTGGTGGCACTTGGCT	121						
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QY	241	TTGCA	245						
DB	60	TTGCA	56						

Search completed: October 24, 2003, 22:51:13
 Job time : 2895.5 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 24, 2003, 15:43:05 : Search time 1284.09 seconds
(without alignments):
:4809 :12 Million cell updates/sec

Title: US-09-830-902-1_COPY_10000_10500
Perfect score: 501
Sequence: 1 ggttcctgtggctcttgggggctgggggagggggcggc 501

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.2

Searched: 2888711 seqs, 2645461386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 3

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*

1: gb_ba.*

2: gb_brg.*

3: gb_brn.*

4: gb_cm.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pr.*

10: gb_rc.*

11: gb_sts.*

12: gb_sy.*

13: gb_un.*

14: gb_vt.*

15: em_ba.*

16: em_fun.*

17: em_hum.*

18: em_in.*

19: em_mu.*

20: em_or.*

21: em_ov.*

22: em_pat.*

23: em_ph.*

24: em_pl.*

25: em_rc.*

26: em_sts.*

27: em_un.*

28: em_vt.*

29: em_hg_hum.*

30: em_hg_inv.*

31: em_hg_inv.*

32: em_hg_inv.*

33: em_hg_inv.*

34: em_hg_inv.*

35: em_hg_inv.*

36: em_hg_inv.*

37: em_hg_inv.*

38: em_hg_inv.*

39: em_hg_inv.*

40: em_hg_inv.*

41: em_hg_inv.*

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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1	501	100.0	101584	9	CNS01255	AL121655 BAC seque
2	501	100.0	110000	6	AX93471	AX93471 Sequence
3	501	100.0	110000	9	HS246003	AX246003 Homo sapi
4	501	100.0	155943	9	AX012364	AC012364 Homo sapi
5	472	94.2	3263	6	AX93472	AX93472 Sequence
6	472	94.2	3263	9	HS246001	AX246001 Homo sapi
7	472	94.2	5120	9	AB229006	AB022906 Homo sapi
8	442	88.2	185281	2	AC112132	AC011232 Homo sapi
9	348.8	69.6	2152	6	AX93576	AX93576 Sequence
10	348.8	69.6	2152	9	HS246003	AX246003 Homo sapi
11	346	69.1	2116	10	BC046286	BC046286 Mus muscu
12	274	54.7	71793	2	AC101733	AC101733 Mus muscu
13	247.4	49.4	721	9	HS246001	AX246001 Homo sapi
14	213.4	42.6	712	9	HS246003	AX246003 Homo sapi
15	165.8	33.1	697	9	HS246003	AX246003 Homo sapi
16	73	14.6	1889	6	AX93542	AX93542 Sequence
17	73	14.6	1889	10	MM1246002	AJ246002 Mus muscu
18	64	12.8	125020	9	AF429315	AF429315 Homo sapi
19	59.4	11.9	161822	2	AC144461	AC144461 Homo sapi
20	58.6	11.7	133575	8	CNS0209N	AC130259 Mus muscu
21	57.2	11.6	37159	2	AC100259	AC100259 Mus muscu
22	57.2	11.4	65942	2	AC031612	AC031612 Homo sapi
23	57	11.4	200542	9	CNS01RG3	AL157736 Human chr
24	56.6	11.3	7218	6	166494	166494 Sequence 14
25	55.8	11.1	135119	2	AC011578	AC011578 Homo sapi
26	55.4	12.1	154729	9	AC018730	AC018730 Homo sapi
27	54.8	10.9	131632	2	AC034263	AC034263 Homo sapi
28	54.4	10.9	80133	2	AC021347	AC021347 Homo sapi
29	54.2	10.8	63082	2	AC022653	AC022653 Homo sapi
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31	53.4	10.7	30693	2	AC079431	AC079431 Mus muscu
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34	53	10.6	159851	2	AP033571	AP033571 Oryza sat
35	53	10.6	171574	2	AC012300	AC012300 Homo sapi
36	53	10.6	219952	2	AC084824	AC084804 Mus muscu
37	52.8	10.5	26822	9	AC009127	AC009107 Homo sapi
38	52.6	10.5	43343	2	AC099831	AC099831 Homo sapi
39	52.4	10.5	2837	9	HSC0G1575	AJ001214 Homo sapi
40	52.4	10.5	139057	2	AP004737	AP004737 Oryza sat
41	52.4	10.5	141534	2	AP033685	AP033685 Oryza sat
42	52.4	10.5	155419	2	AP058403	AP058403 Oryza sat
43	52.4	10.5	180865	2	AP033766	AP033766 Oryza sat
44	52.2	10.4	64211	2	AC091110	AC091110 Homo sapi
45	52.2	10.4	133790	2	AC119292	AC119292 Oryza sat

ALIGNMENTS

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LOCUS
DEFINITION
BAC sequence from the SP04 candidate region at 2p21-2p22 BAC 316p14
of library CITR_978_SKB from chromosome 2 of Homo sapiens (Human).
ACCESSION
AL121655
VERSION
AL121655.1
KEYWORDS
SP04 genomic DNA interval.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1. Bases 1 to 101584;
Hazar, C., Ponkrecht, N., Mavel, D., Farnet, C., Samson, D.,
Attiguenave, F., Davoine, C., Cruaud, C., Durr, A., Wincker, P.,

Pred. No. is the number of results predicted by chance to have a


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Best Local Similarity 100.0%; Pred. No. 1,4e-74;
Matches 501; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY : GGTTCGGTCTCGGGAGCGGGTATGCGCGCGCGGGCAGTGAGAGCTGTGAATG 60
Db 10000 GGTTCGGTCTCGGTGCGGAGCGGGTATGCGCGCGCGGGCAGTGAGAGCTGTGAATG 10059
QY 6: AATTCGGGTGAGCAGGAGAGAGAGAGGCTCGCGCGCGCGCGCGCGCGCTGCCT 120
Db 10060 AATTCGGGTGAGCAGGAGAGAGAGAGGCTCGCGCGCGCGCGCGCGCGCTGCCT 10119
QY 121 CCCAGGCTCCGGCCCTTGCCTGGCCCGCCCGCCCTCCGCGCGCGCGCGCGCGCTCCG 180

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Db 10120 CCCAGGCTCCGGCCCTTGCCTGGCCCGCCCGCCCTCCGCGCGCGCGCGCTCCG 10179
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Db 10180 CCCAGTTCGGGATAGCGGAACCTGTACTATTTCTCTAGCCGCTGTTGTAGCTTC 10239
QY 241 GCGCTGCTGGTTTGGTTCGCTTCGACCTGGGCTCTCTTCTGCTGCTGCGCAGGC 300
Db 10240 GCGCTGCTGGTTTGGTTCGCTTCGACCTGGGCTCTCTTCTGCTGCTGCGCAGGC 10299
QY 301 TTCTCCGCGCCCTCATGGCAGCAAGAGAGCTCGGGGCGCGCGCGCAGCAGCTGCTCG 360
Db 10300 TTCTCCGCGCCCTCATGGCAGCAAGAGAGCTCGGGGCGCGCGCGCAGCAGCTGCTCG 10359
QY 361 GCGTCCGCGCGCGCGCGGCTCGGGGCGCGGAGCGCGAGCGCTCGAGTCTTCCACAA 420
Db 10360 GCGTCCGCGCGCGCGCGGCTCGGGGCGCGGAGCGCGAGCGCTCGAGTCTTCCACAA 10419
QY 421 CAGGCTTCGAGTACATCTCCATTCGCTGCGCATGATGAGATGAGAAAGGTAAGTAG 480
Db 10420 CAGGCTTCGAGTACATCTCCATTCGCTGCGCATGATGAGATGAGAAAGGTAAGTAG 10479
QY 481 GGGCTGGGGGAGGGGCGGC 501
Db 10480 GGGCTGGGGGAGGGGCGGC 10500

RESULT 4
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LOCUS Homo sapiens BAC clone RP11-444D15 from 2, complete sequence.
DEFINITION AC012364
ACCESSION AC012364
VERSION AC012364.7 GI:14589736
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 155943)
AUTHORS Salton, J.E. and Waterston, R.
TITLE Toward a complete human genome sequence
JOURNAL Genome Res. 8 (11), 1097-1109 (1998).
MEDLINE 98263792
PMID 9847074
REFERENCE 2 (bases 1 to 155943)
AUTHORS Ali, N. and Abbott, A.
TITLE The sequence of Homo sapiens BAC clone RP11-444D15
JOURNAL Unpublished (2001)
REFERENCE 3 (bases 1 to 155943)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (25-Oct-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE 4 (bases 1 to 155943)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (03-Jul-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE 5 (bases 1 to 155943)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (09-JAN-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Jul 3, 2001 this sequence version replaced gi:1162953.
COMMENT
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@wustl.wustl.edu
----- Summary Statistics
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Center project name: H_NH3244D15

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NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressors and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RPL11 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Woon, P.-Y., Zhao, B., Frongier, E., Rateno, M., Catanesse, J. J., and de Jong, P. J. (1999). An improved approach for construction of bacterial artificial chromosome libraries. *Genetics* 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu/>).

VECTOR: pBAC3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RPL1-41E19; the clone sequenced to the right is RPL1-26E13. Actual start of this clone is at base position 1 of RPL1-444D15; actual end is at base position 155943 of RPL1-444D15.

Data from AC011232 and AC010961 was used to finish this clone, AC012364. The sequence fidelity between bases 112626 to 112643 can not be guaranteed due to an unresolved homopolymeric run. The sequence between 112618 to 112663 is single stranded.

FEATURES

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DB 125853 CGGAGGCTCCGCGGCTTCCCTGCGGCGCGGCGGCTCCGCGGCGCGGCGGCTCCG 125912
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DEFINITION Sequence 2 from Patent WO0119198.
ACCESSION AX093472
VERSION AX093472.1 G1:135099:2
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE
AUTHORS Hazan, J., Farkasch, N., Mavel, D., Paternotte, C., Samson, D.,
Aitguenave, F., Davoine, C., Cruaud, C., Durr, A., Winkler, P.,
Bottier, P., Cattolico, L., Barbe, V., Burgunder, J.M.,
Frid'homme, G.F., Brice, A., Fontaine, B., Heilig, R. and
Weissenbach, J.
TITLE Spastin, a new AAA protein, is altered in the most frequent form of
autosomal dominant spastic paraplegia
JOURNAL Nat. Genet. 23 (3), 296-303 (1999)
MEDLINE 20055425
PUBMED 10610178
REFERENCE
AUTHORS Farkasch, N., Mavel, D., Byrne, P., Davoine, C., Cruaud, C.,
Beetsch, D., Samson, D., Coutinho, P., Hutchinson, M., McMoragie, P.,
Burgunder, J., Tartaglione, A., Heinzel, C., Fekli, I., Deufel, T.,
Pafrey, N., Brice, A., Fontaine, B., Prud'homme, J., Weissenbach, J.,
Durr, A. and Hazan, J.
TITLE Spectrum of SPG4 mutations in autosomal dominant spastic paraplegia
JOURNAL Hum. Mol. Genet. 9 (4), 637-644 (2000)
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MEDLINE 20164302
PUBMED 10699187
REFERENCE 3 (bases 1 to 3263):
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (17-JUN-1999) Genoscope, Genoscope - Centre National de
Sequencage, BP 191, EVRY 91006, FRANCE
COMMENT E-mail : seqref@genoscope.cns.fr - Web : www.genoscope.cns.fr.
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69 GGTTCGGTTCGGTCTGCGGAGCGGGTATGCGGCGCGGCGGAGTGTGAATG 129
QY |||||
61 AATTCTCCGGTGAAGCGGAAAGAAAGGCTCGGCGCGGCGGCGGAGTGTGAATG 120
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QY |||||
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DQ |||||
189 CCCAGGCTCGGCGGCTTCGCTGGCGCGGCGGCGGCTCGGCGCGGCGGCGGCTCG 248
QY |||||
181 CCCAGTTCGGCGGCTAAGCGGAAAGCTGTACTATTTCTCTACCGGCTGTGTGAATG 240
DQ |||||
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DQ |||||
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421 CAGGCTTCGAGTACATCTCTGATTCGCTTCGCTTCGCTTCGCTTCGCTTCGCTTC 492
DQ |||||

DQ 489 CAGGCTTCGAGTACATCTCTGATTCGCTTCGCTTCGCTTCGCTTCGCTTCGCTTC 540
RESULT 7
LOCUS 5120 bp mRNA linear PRI 04-AUG-1999
DEFINITION Homo sapiens mRNA for KIAA1063 protein, complete cds.
ACCESSION AB029006
VERSION AB029006.1 GI:5689502
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
2 (sites)
Xikuro.R., Nagase.T., Ishikawa.K., Hirosewa.M., Miyajima.N.,
Tanaka.A., Kotani.H., Nomura.N. and Chara.C.
Prediction of the coding sequences of unidentified human genes.
XIV. The complete sequences of 100 new cDNA clones from brain which
code for large proteins in vitro
cDNA Res. 6 (3), 197-205 (1999)
JOURNAL
MEDLINE 95397452
PUBMED 10470851
REFERENCE 2 (bases 1 to 5120)
AUTHORS Chara, O., Nagase, T. and Kikuno, R.
TITLE Direct Submission
JOURNAL Submitted (17-JUN-1999) Osamu Ohara, Kazusa DNA Research Institute,
Laboratory of DNA Technology, Yana 1532-3, Kisarazu, Chiba
212-0912, Japan (E-mail:cdna@kazusa.or.jp, Tel.:81-438-52-3913,
Fax:81-438-52-3914)
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DB 23262 GGGGCTGGGAGGGGGGGC 23283

RESULT 9
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DEFINITION Sequence 126 from Patent WO0118198.
ACCESSION AX091576
VERSION AX091576.1 GI:13510014
KEYWORDS Mus musculus (house mouse);
SOURCE Mus musculus
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1
AUTHORS Weissenbach, J. and Hatan, J.
TITLE Cloning, expression and characterisation of the spg4 gene
responsible for the most frequent form of autosomal spastic
paraplegia
JOURNAL Patent: WO 0118198-A 106 15-MAR-2001;
CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (CNRS) (FR)
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Best Local Similarity 86.7%; Fred. No. 7.3e-49;
Matches 409; Conservative 0; Mismatches 57; Indels 6; Gaps 2;
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DB 493 CAGGCGCTTCAGTACATCTCCATTCGCTGGCAGTCAGAGAGAGAGAGAGAGAG 544

RESULT 11
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LOCUS 2116 bp mRNA linear ROD 10-FEB-2003
DEFINITION Mus musculus, spastic paraplegia 4 homo-log (human), clone MSC:54786
IMAE:6441742, mRNA, complete cds.
ACCESSION BC046286
VERSION BC046286.1 GI:28279481

ACCESSION NB6-679R.
VERSION AJ325510
KEYWORDS GAG325510.1 GI:13869904
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 969)
AUTHORS Kutsenko, A.S., Gazatulin, R.Z., Al-Amin, A.N., Wang, F., Kvasha, S.M.,
Podolski, R.M., Matushkin, Y.G., Gvanchandani, A., Muravenko, O.V.,
Levitsky, V.G., Kolesnikov, N.A., Protopopov, A.I., Kasha, V.I.,
Kisselev, J.L., Wasserman, W., Wahlstedt, C.I. and Zharovskiy, E.R.
TITLE Not1 flanking sequences: a tool for gene discovery and verification
of the human genome
JOURNAL Nucleic Acids Res. 30 (14): 3153-3170 (2002)
MEDLINE 2213767
PUBMED 12136098
REFERENCE 2 (bases 1 to 969)
AUTHORS Zharovskiy, E.R.
TITLE Direct Submission
JOURNAL Submitted (16-MAY-2001) Microbiology and Tumorbiology Centre,
Karolinska Institute, Theorell's vag, 3, Box 280, Stockholm 171 77,
Sweden
FEATURES
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Best Local Similarity 92.8%; Fred. No. 1.1e-48;
Matches 376; Conservative 0; Mismatches 28; Indels 1; Gaps 1;
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DB 220 GGTTCGCTGGTCTGGGAGGGGGTATGGGGGGGGGGAGAGAGTGTGAATG 279
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DB 280 AATTTCGGGTGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 339
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QY 181 CCGAGTCCCGCATAAGCGGAAGTGTACTATTCTCTACCGCTGTTGTAGGCTTC 240
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QY 241 GGGCTGTGGGTTGGTGGCTTCACCTTGGGCTGCTTCTGGTGGCTTGGCAGGC 300
DB 460 GGGCTGTGGGTTGGTGGCTTGGCAGTGGGCTTGGCAGTGGGCTTGGCAGGC 519
QY 301 TTCTCCCGCGCCCTATGCGAGCGAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 360
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QY 361 GCGTGGGCGCGGCGGCTGGCGGCGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 405
DB 580 GCGTGGGCGCTCGCGCAGTTCGCGGCGG-GAGGCGGAACGCGTC 623

RESULT 11
PC046286
LOCUS 2116 bp mRNA linear ROD 10-FEB-2003
DEFINITION Mus musculus, spastic paraplegia 4 homo-log (human), clone MSC:54786
IMAE:6441742, mRNA, complete cds.
ACCESSION BC046286
VERSION BC046286.1 GI:28279481
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5

TITLE
JOURNAL
COMMENT

Direct Submission
Submitted (23-NOV-2003) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
All repeats were identified using RepeatMasker:
Sait, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
..... Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIGR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence.submissions@genome.wi.mit.edu
----- Project Information
Center project name: 31731
Center clone name: 4.7.1_6

* NOTE: This record contains 91 individuals.
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.

591	690:	contig of 690 bp in length	18953	19056:	gap of 100 bp
791	790:	gap of 100 bp	19059	19757:	contig of 699 bp in length
1465	1465:	contig of 675 bp in length	19758	19857:	gap of 100 bp
1565	1565:	gap of 100 bp	19858	20513:	contig of 653 bp in length
2271	2271:	contig of 705 bp in length	20511	20610:	gap of 100 bp
2371	2371:	gap of 100 bp	20611	21295:	contig of 685 bp in length
3056	3056:	contig of 695 bp in length	21296	21395:	gap of 100 bp
3166	3166:	gap of 100 bp	21396	22087:	contig of 692 bp in length
3651	3651:	contig of 685 bp in length	22088	22187:	gap of 100 bp
3852	3852:	gap of 100 bp	22188	22860:	contig of 673 bp in length
4337	4337:	contig of 685 bp in length	22861	23646:	contig of 666 bp in length
4737	4737:	gap of 100 bp	23647	23746:	gap of 100 bp
5436	5436:	gap of 100 bp	23747	24439:	contig of 683 bp in length
5537	5537:	contig of 698 bp in length	24440	24529:	gap of 100 bp
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7134	7134:	contig of 688 bp in length	25999	26098:	gap of 100 bp
7821	7821:	gap of 100 bp	26100	26800:	contig of 702 bp in length
7822	7822:	contig of 689 bp in length	26801	26900:	gap of 100 bp
8611	8611:	gap of 100 bp	26901	27613:	contig of 719 bp in length
8711	8711:	contig of 701 bp in length	27620	28436:	contig of 719 bp in length
9412	9412:	contig of 701 bp in length	28437	28536:	gap of 100 bp
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			46578	46577:	gap of 100 bp
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			47363	47462:	gap of 100 bp

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12136098
REFERENCE 2 (bases 1 to 721):
AUTHORS Zabarovsky,E.R.
TITLE Direct Submission
JOURNAL Submitted (16-MAY-2001) Microbiology and Tumourbiology Centre,
Karolinska Institute, Theorells vag, 3, Box 280, Stockholm 171 77,
Sweden
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Best Local Similarity 95.0%; Pred. No. 7 4e-32;
Matches 266; Conservative 0; Mismatches 12; Indels 2; Gaps 1
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DEFINITION HSJ-6A3ARS.
ACCESSION  AF342199
VERSION     AF342199
KEYWORDS   AF342199.1  GI:15986616
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
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            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 712):
AUTHORS     Kurzenko,A.S., Gizatullin,R.Z., Al-Amin,A.N., Wang,F., Kvasha,S.M.,
            Podolski,R.M., Katushkin,Y.G., Gyarachandani,A., Muravenko,O.V.,
            Lovitskiy,V.G., Kolchanov,N.A., Protopopov,A.I., Kashiba,V.I.,
            Kasselev,I.B., Wasserman,W., Wahlstedt,C. and Zabarovsky,E.R.
            NstI flanking sequences: a tool for gene discovery and verification:
            of the human genome
JOURNAL     Nucleic Acids Res. 30 (14), 3161-3170 (2002)
MEDLINE     2213767
REFERENCE   2 (bases 1 to 712)
AUTHORS     Zabarovsky,E.R.
TITLE       Direct Submission
JOURNAL     Submitted (16-MAY-2001) Microbiology and Tumourbiology Centre,
            Karolinska Institute, Theorells vag, 3, Box 280, Stockholm 171 77,
            Sweden
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XX FR2798138-A1.
XX 09-MAR-2001.
XX 03-SEP-1999; 99PR-0011097.
XX 03-SEP-1999; 99PR-0011097.
XX CNRS ; CNRS CENT NAT RECH SCI.
XX Weissenbach J, Hazan J;
XX WPI; 2001-283966/30.
XX New human nucleic acid from the SPG4 gene, useful e.g. for diagnosis of
XX autosomal dominant familial spastic paraplegia and in drug screening .
XX Claim 2; Page 45-106; 145pp; French.
XX The present sequence represents a human SPG4 gene. The SPG4 gene encodes
XX a spastin polypeptide. Mutations in the SPG4 gene are responsible for
XX autosomal dominant familial spastic paraplegia. SPG4 polynucleotides,
XX and their fragments, are used to screen DNA banks for sequences that
XX encode spastin (particularly sequences in other mammals, specifically
XX mice); to identify SPG4 mutations, or other genetic anomalies.
XX particularly for diagnosis of autosomal dominant familial spastic
XX paraplegia (PSP-AD); to identify promoters and other regulatory elements
XX of the SPG4 gene; for detection and amplification; for recombinant
XX production of spastin; and for diagnostic genotyping of PSP-AD.
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XX Best Local Similarity 100.0%; Pred. No. 8.7e-92;
XX Matches 501; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB 10000 GGTTCGGTGGTGTGCGGAGCGGGGTATGCGGCGGCGGCGGAGTGAGAGCTGTGATG 10000
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DB 10240 TTCTCCGCGCGCTCTCATGGCAGGCAAGAGAGTCTCCGGGCGGCGGCGGAGACCTGCTCG 363

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CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one
CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
CC and the degree of hybridisation to both classes is determined from the
CC label on the amplicon. From the ratio of labels hybridised to the two
CC classes of oligomers, the degree of methylation is calculated. The method
CC is used: (1) for diagnosis and/or prognosis of side effects of
CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
CC of the central nervous, cardiovascular, gastrointestinal and respiratory
CC systems etc., particularly by detecting mutations or single nucleotide
CC polymorphisms (SNPs); and (2) for differentiation of cell or tissue
CC types and for investigating cell differentiation. The method allows the
CC methylation status of many C residues to be determined simultaneously.
CC AB013410-AB054121 represent genomic DNA sequences used to illustrate the
CC method for determining the degree of cytosine methylation described in
CC the disclosure of the invention.

SQ Sequence 777 BP; 135 A; 79 C; 261 G; 302 T; 0 Other;

Query Match: 63.0%; Score 315.4; DB 24; Length 777;
Best Local Similarity 78.8%; Pred. No. 1.4e-54;
Matches 376; Conservative 0; Mismatches 101; Indels 0; Gaps 0;
QY 3 TTCCCGTGGTCTGGGAGCGGGTTATGGCGCGCGGCGGAGTGGAGCTGTGATGAA 62
DB 668 TTCCCGTGGTCTGGGAGCGGGTTATGGCGCGCGGCGGAGTGGAGCTGTGATGAA 609
QY 63 TTCTCCGGTGGAGCGGGAAGAAAGCTCCGGCGCGGCGGAGTGGAGCTGTGATGAA 122
DB 608 TTCTCCGGTGGAGCGGGAAGAAAGCTCCGGCGCGGCGGAGTGGAGCTGTGATGAA 549
QY 123 CAGGCGTCCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 182
DB 545 CAAACCTCCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 489
QY 183 CGAGTCCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 242
DB 488 CGAATCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 429
QY 243 GGTGCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 302
DB 428 GCTACTACGTTTAACTGGCTTCCACCTAAACCTCTCTGATATACTCTACCAAGCTT 369
QY 303 CTGCGCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 362
DB 369 CTGCGCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 309
QY 363 CTGCG 422
DB 308 CTGCG 249
QY 423 GGTGCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 479
DB 248 AACTCTGGATATATCTCCATTCCTACCGATCGATATAAATAAATAAATAAATAA 192

RESULT 4

AB039713

10 AB039713 standard; DNA; 777 BP.

XX AB039713;

XX AC

XX XX

DC 12-JUL-2002 (first entry);

XX

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 26124.

XX

KW Human; cytosine methylation; 5'-CpG-3', uracil; cytosine; diagnosis;

KW drug; side effect; cancer; central nervous system; cardiovascular;

KW gastrointestinal; respiratory system; single nucleotide polymorphism;

KW SNP; cell differentiation; ds.

XX

OS Homo sapiens.

XX WO200228632-A2.
PN
XX
XX PC
XX 07-MAR-2002.
XX
XX 01-SEP-2002; 2001WO-EF10074.
XX
XX 01-SEP-2002; 2000DE-1043826.
XX 05-SEP-2000; 2000DE-1044543.
XX (EPIG-) EPIGENOMICS AG.
XX
XX Ciek A, Piepenbrock C, Berlin K, Gueling D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful
XX for diagnosis and prognosis, comprises selective hybridization of
XX amplicons from chemically treated DNA -
XX
XX Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
XX methylation of a particular cytosine in a motif 5'-CpG-3', present in a
XX genomic sample of DNA. The sample is treated chemically to convert
XX cytosine (C) but not methylated C, to uracil, then part of the genomic
XX DNA that contains the target C is amplified to form a labeled amplicon.
XX The amplicon is hybridised to two classes, each with at least one
XX member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
XX and the degree of hybridisation to both classes is determined from the
XX label on the amplicon. From the ratio of labels hybridised to the two
XX classes of oligomers, the degree of methylation is calculated. The method
XX is used: (1) for diagnosis and/or prognosis of side effects of
XX therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
XX of the central nervous, cardiovascular, gastrointestinal and respiratory
XX systems etc., particularly by detecting mutations or single nucleotide
XX polymorphisms (SNPs); and (2) for differentiation of cell or tissue
XX types and for investigating cell differentiation. The method allows the
XX methylation status of many C residues to be determined simultaneously.
XX AB013410-AB054121 represent genomic DNA sequences used to illustrate the
XX method for determining the degree of cytosine methylation described in
XX the disclosure of the invention.

SQ Sequence 777 BP; 302 A; 261 C; 79 G; 135 T; 0 Other;

Query Match: 63.0%; Score 315.4; DB 24; Length 777;

Best Local Similarity 78.8%; Pred. No. 1.4e-54;

Matches 376; Conservative 0; Mismatches 101; Indels 0; Gaps 0;

QY 3 TTCCCGTGGTCTGGGAGCGGGTTATGGCGCGCGGCGGAGTGGAGCTGTGATGAA 62
DB 110 TTCCCGTGGTCTGGGAGCGGGTTATGGCGCGCGGCGGAGTGGAGCTGTGATGAA 169
QY 63 TTCTCCGGTGGAGCGGGAAGAAAGCTCCGGCGCGGCGGAGTGGAGCTGTGATGAA 122
DB 120 TTCTCCGGTGGAGCGGGAAGAAAGCTCCGGCGCGGCGGAGTGGAGCTGTGATGAA 229
QY 123 CAGGCGTCCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 182
DB 230 CAAACCTCCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 269
QY 183 CGAGTCCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 242
DB 290 CGAATCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 349
QY 243 GGTGCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 302
DB 250 GCTACTACGTTTAACTGGCTTCCACCTAAACCTCTCTGATATACTCTACCAAGCTT 479
QY 303 CTGCGCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 362
DB 410 CTGCGCGCGCGCTTGGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 469

XX PS Claim 12: 56pp + Sequence Listing: 56pp: German.

XX This invention describes a novel method for determining the degree of methylation of a particular cytosine in a motif 5'-CpG-3', present in a genomic sample of DNA. The sample is treated chemically to convert cytosine (C) but not methylated C to uracil, then part of the genomic DNA that contains the target C is amplified to form a labeled amplicon. The amplicon is hybridised to two classes, each with at least one member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the degree of hybridisation to both classes is determined from the label on the amplicon. From the ratio of labels hybridised to the two classes of oligomers, the degree of methylation is calculated. The method is used: (i) for diagnosis and/or prognosis of side effects of therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders of the central nervous, cardiovascular, gastrointestinal and respiratory systems etc., particularly by detecting mutations of single nucleotide polymorphisms (SNP's); and (ii) for differentiation of cell or tissue types and for investigating cell differentiation. The method allows the methylation status of many C residues to be determined simultaneously. ABQ3410-ABQ54121 represent genomic DNA sequences used to illustrate the method for determining the degree of cytosine methylation described in the disclosure of the invention.

XX SQ Sequence 777 BP: 317 A; 275 C; 79 G; 106 T; 0 other;

Query Match: 60.4%; Score 302.6; DB 24; Length 777;
 Best Local Similarity 75.2%; Pred. No. 5.3e-52;
 Matches 377; Conservative C; Mismatches 124; Indels 0; Gaps 0;

QY 1 GGTTCCTCGCTGCTGGGAGCGGGTATGCGCGCGGCGCGCTGAGAGCTGTGATG 60
 DB 670 GGTTCCTCGCTGCTGGGAGCGGGTATGCGCGCGGCGCGCTGAGAGCTGTGATG 611

QY 61 AATTCCTCGGCTGGAGCGGGAAGAAAGAGCTCCGCGCGGCGCGAGCAACCGGTGCT 120
 DB 610 AATTCCTCGGCTGGAGCGGGAAGAAAGAGCTTCGCGCGGCGGTAGTAATCGGTGT 551

QY 121 CCGAGGCTCCGCGGCTGCTGCGCGCGCGCGCGCTTCGCGCGGCGCGCGCGCTCG 180
 DB 550 TTTAGGTTTCGTTTGTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTT 491

QY 181 CCGAGTCCCGCATAGCGGAGCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 240
 DB 490 TCGAGTCCGCTATAGCGGAGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTT 431

QY 241 GCGTCTCGGTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 300
 DB 430 GCGTCTCGGTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 371

QY 301 TCTCCGCGCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 360
 DB 370 TTTTTCGCGTTTATATGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGCT 311

QY 361 GCTTCGCGCGCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 420
 DB 310 GTTCGTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 251

QY 421 CAGGCTTCGTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 480
 DB 250 TAGGTTTCGTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 191

QY 481 GGGGTGGGGAGGGGGGGG 501
 DB 190 GGGGTGGGGAGGGGGGGG 170

RESULT 7
 ABQ24124/c
 ID ABQ24124 standard; DNA: 562 BP.
 XX
 AC ABQ24124;
 XX

DT 12-JUL-2002 (first entry);

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO: 10715.

XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
 KW drug; side effect; cancer; central nervous system; cardiovascular;
 KW cytosine (C) but not methylated C; to uracil; then part of the genomic
 KW gastrointestinal; respiratory system; single nucleotide polymorphisms;
 KW SNP; cell differentiation; ds.
 XX Homo sapiens.
 OS
 XX KW2002:5632-A2.
 PN 07-MAR-2002.
 FD
 XX 01-SEP-2001: 2001WO-EP10074.
 XX 01-SEP-2000: 2000DS-1543826.
 XX 05-SEP-2000: 2000DS-1044543.
 XX (EPIG-) BPGENOMICS AG.
 PA
 XX Olek A. Piepenhock C, Berlin K. Guetig D;
 PI
 XX WPI: 2002-371829/40.
 DR
 XX Determining the degree of cytosine methylation in genomic DNA, useful
 PI for diagnosis and prognosis, comprises selective hybridization of
 PT amplicons from chemically treated DNA .
 XX
 XX Claim 12: 56pp - Sequence Listing: 56pp: German.

XX This invention describes a novel method for determining the degree of methylation of a particular cytosine in a motif 5'-CpG-3', present in a genomic sample of DNA. The sample is treated chemically to convert cytosine (C) but not methylated C to uracil, then part of the genomic DNA that contains the target C is amplified to form a labeled amplicon. The amplicon is hybridised to two classes, each with at least one member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the degree of hybridisation to both classes is determined from the label on the amplicon. From the ratio of labels hybridised to the two classes of oligomers, the degree of methylation is calculated. The method is used: (i) for diagnosis and/or prognosis of side effects of therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders of the central nervous, cardiovascular, gastrointestinal and respiratory systems etc., particularly by detecting mutations of single nucleotide polymorphisms (SNP's); and (ii) for differentiation of cell or tissue types and for investigating cell differentiation. The method allows the methylation status of many C residues to be determined simultaneously. ABQ3410-ABQ54121 represent genomic DNA sequences used to illustrate the method for determining the degree of cytosine methylation described in the disclosure of the invention.

XX SQ Sequence 562 BP: 113 A; 53 C; 195 G; 201 T; 0 other;

Query Match: 45.9%; Score 229.6; DB 24; Length 562;
 Best Local Similarity 50.3%; Pred. No. 2.2e-37;
 Matches 282; Conservative C; Mismatches 67; Indels 2; Gaps 1;

QY 129 TCGCGCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 188
 DB 562 TCGCGCGCTTACCTAACCGCGCGCTTCCGCGCGCGGAGACACCTCCGCGCGAATC 503

QY 189 GCGCATAGCGGAAGCTGTACTATTTCTCTACCGCGCTGTTGTAGGCTTCGCGTGT 248
 DB 502 GCGCATAGCGGAAGCTGTACTATTTCTCTACCGCGCTTATATAAAGCTTCGCGTGT 443

QY 249 GCGTTTGGTTCGCTTCCACTGCGGCTTCCTCTTCGTGTGGCTTCGCGAGGCTTTCCCG 308
 DB 442 ACGTTTAAATCGCTTCGACTAAAGCTTCCTCTTCGTATTAACCTTACCAAGGCTTCTCCG 383

QY 309 CGCCTCATGGCAGCAAGAGGAGCTCCGCGCGCGCGGAGGAGCTGCTTCGCGCTCGG 368

DB 382 CCCCCTCTATACACCAAAACAACTCCGAAACCCGGCGGACACACTACCTCGACCTCGAC 323
QY 369 CCGCGCGCGGTGCGCGGCGGAGCGGCGGTTCGAGTCTTCACAGAAACAGGCTT 428
DB 322 CC--GACGCCATACCGACGACGAAACCGAAGCGTTCGGAATCTTCACAAACAACTT 265
QY 429 CGAGTACATCTCCATTCGCTCGGCGATCGATGAGGATGAGAAAGGTAACTA 479
DB 264 CGAATACATCTCCATTCGCTCGGCGATCGATGAGGATGAGAAAGGTAACTA 214

RESULT 8
ABQ24:125
ID ABQ24:125 standard; DNA; 562 BP.
XX
AC ABQ24:125;
XX
CT 12-JUL-2002 (first entry);
XX
DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 10713.
XX
KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX Gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.
XX
OS Homo sapiens.
XX
FN WC200218632-A2.
XX
PD 07-MAR-2002.
XX
PE 01-SEP-2000; 2001WO-EP10074.
XX
PR 01-SEP-2000; 2000DE-1043826.
XX
PR 05-SEP-2000; 2000DE-1044543.
XX
PA (EP1G-) EP1GENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K, Gueting D;
XX
DR WPI; 2002-371829/40.
XX

PT Determining the degree of cytosine methylation in genomic DNA, useful
PT for diagnosis and prognosis, comprises selective hybridization of
PT amplicons from chemically treated DNA
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridized to two classes, each with at least one
CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
CC and the degree of hybridization to both classes is determined from the
CC label on the amplicon. From the ratio of labels hybridized to the two
CC classes of oligomers, the degree of methylation is calculated. The method
CC is used: (i) for diagnosis and/or prognosis of side effects of
CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
CC of the central nervous, cardiovascular, gastrointestinal and respiratory
CC systems etc., particularly by detecting mutations or single nucleotide
CC polymorphisms (SNPs); and (ii) for differentiation of cell or tissue
CC types and for investigating cell differentiation. The method allows the
CC methylation status of many C residues to be determined simultaneously.
CC ABQ14:0-ABQ54:121 represent genomic DNA sequences used to illustrate the
CC method for determining the degree of cytosine methylation described in
CC the disclosure of the invention.
XX
SQ Sequence 562 BP; 201 A; 195 C; 53 G; 113 T; 0 other;

Query Match 45.9%; Score 229.8; DS 24; Length 582;

Best Local Similarity 80.3%; Pred. No. 2,2e-37;
Matches 282; Conservative C; Mismatches 67; Indels 2; Gaps 1;
QY 129 TCCGCCCCCTTGGCTGGCGCCCGCCCTCCGCGCGCGGGCGCGCCCTCGCGCGGAGTC 188
DB TCCGCCCCCTTACCTAACCCCGCCCTCCGCGCGCGGAAACCGACCCCTCGCGCGGATC 60
QY 189 CCGCGATTAAGCGGAACCTGTACTATTCTCTACCCCTGTTTCTAGGCTTCGCGTCT 248
DB 61 CCGCGATAAAGCAAACTATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 120
QY 249 CCGTTTGGTGGCTTCCACCTGGGGCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 328
DB 121 AGTTTAAATCGCTTCCACCTAAACCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 180
QY 389 CCGCTCATGCGACCAAGAGAGGCTCGGGCGCGCGCGCGCGCGCGCGCGCGCGCG 368
DB 181 CCGCTCATGCGACCAAGAGAGGCTCGGGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 240
QY 369 CCGCGCGCGGTGCG 428
DB 241 CC--GACGCCATACCGACGACGAAACCGAAGCGTTCGGAATCTTCACAAACAACTT 298
QY 429 CGAGTACATCTCCATTCGCTCGGCGATCGATGAGGATGAGAAAGGTAACTA 479
DB 264 CGAATACATCTCCATTCGCTCGGCGATCGATGAGGATGAGAAAGGTAACTA 214

RESULT 9
ABQ24:122
ID ABQ24:122 standard; DNA; 562 BP.
XX
AC ABQ24:122;
XX
CT 12-JUL-2002 (first entry)
XX
DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 10713.
XX
KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX Gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.
XX
OS Homo sapiens.
XX
FN WC200218632-A2.
XX
PD 07-MAR-2002.
XX
PE 01-SEP-2000; 2001WO-EP10074.
XX
PR 01-SEP-2000; 2000DE-1043826.
XX
PR 05-SEP-2000; 2000DE-1044543.
XX
PA (EP1G-) EP1GENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K, Gueting D;
XX
DR WPI; 2002-371829/40.
XX

PT Determining the degree of cytosine methylation in genomic DNA, useful
PT for diagnosis and prognosis, comprises selective hybridization of
PT amplicons from chemically treated DNA
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.

CC This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridized to two classes, each with at least one
CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers


```

XX
AC AAF84875;
XX
XX 09-JUL-2001 (first entry)
XX
DE Nucleotide sequence of a partial murine SPG4 polypeptide.
XX
KW Mouse: SPG4 gene; spastin; PSP-AD; gene therapy;
KW Autosomal dominant familial spastic paraplegia; ss.
XX
XX
OS
XX
XX Key Location/Qualifiers
XX CDS 3..1517
XX FT /*tag= a
XX FT /product= "spastin"
XX
XX FR2799138-A1.
XX
XX 09-MAR-2001.
XX
XX C3-SEP-1999; 99EP-0011097.
XX
XX C3-SEP-1999; 99EP-0011097.
XX (CNRS) CNRS CENT NAT RECH SCI.
XX
XX Weissenbach J, Hazan J;
XX WPI; 2001-293566/30.
XX P-PSPB; AA668136.
XX
XX New human nucleic acid from the SPG4 gene, useful o.g. for diagnosis of
XX autosomal dominant familial spastic paraplegia and in drug screening.
XX
XX Claim 2; Page 123; 145pp; French.
XX
XX The present sequence encodes a partial spastin polypeptide. Spastin is
XX encoded by the SPG4 gene. Mutations in the SPG4 gene are responsible for
XX autosomal dominant familial spastic paraplegia. SPG4 polynucleotides,
XX and their fragments, are used to screen DNA banks for sequences that
XX encode spastin (particularly sequences in other mammals, specifically
XX mice); to identify SPG4 mutations, or other genetic anomalies.
XX particularly for diagnosis of autosomal dominant familial spastic
XX paraplegia (PSP-AD); to identify promoters and other regulatory elements
XX of the SPG4 gene; for detection and amplification; for recombinant
XX production of spastin; and for diagnostic genotyping of PSP-AD.
XX
XX Sequence 1689 BP; 525 A; 357 C; 416 G; 391 T; 0 other;
XX
XX Query Match 14.6%; Score 73; DB 22; Length 1689;
XX Best Local Similarity 93.8%; Pred. No. 73e-36;
XX Matches 76; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
XX
XX 392 AGCCGAGCGCGTCCGAGCTTTCACAAACAGCCCTTCGAGACATCCCATTCGCCCTGC 451
XX ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX Db 1 AGCCGAGAGCGCGTCCGAGCTTTCACAAACAGCCCTTCGAGACATCCCATTCGCCCTGC 60
XX
XX 452 GCATCGATGGGATGGAAG 472
XX ||||| ||||| ||||| |||||
XX Cc 6 GCATCGAGGAGGAGGAAG 91
XX
XX
XX ABS71875;
XX
XX ABS71875;
XX
XX 02-DEC-2002 (first entry)
XX
XX Human GTP-Rho binding protein 2 single-exon probe #16.
XX
XX

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KW Human; ss; GTP-Rho binding protein 2; GRBP2; chromosome 19q12;
KW oncogene; tumour; liposarcoma; ichthyosis congenita III; probe;
KW benign familial infantile convulsion; gene therapy.
XX
XX Homo sapiens.
XX
XX EP1211216-A2.
XX
XX 14-AUG-2002.
XX
XX 17-JAN-2002; 2002EP-0001C26.
XX
XX 30-JAN-2001; 2001WO-US03663.
XX 30-JAN-2001; 2001WO-US03664.
XX 30-JAN-2001; 2001WO-US03665.
XX 30-JAN-2001; 2001WO-US03666.
XX 30-JAN-2001; 2001WO-US03667.
XX 30-JAN-2001; 2001WO-US03668.
XX 30-JAN-2001; 2001WO-US03669.
XX 30-JAN-2001; 2001WO-US03670.
XX 23-JUN-2001; 2001US-0895040.
XX
XX AECOM-1; AECOMICA INC.
XX
XX Shannon ME, Ji Y;
XX
XX WPI; 2002-694026/74.
XX
XX Novel GTP-Rho binding protein 2 and nucleic acids encoding the protein,
XX useful for the manufacture of a medicament for treating a disease
XX associated with altered expression or activity of human GRBP2 protein.
XX
XX
XX Example 4; Page 57-58; 101pp; English.
XX
XX The invention relates to an isolated GTP-Rho binding protein 2 (GRBP2)
XX polypeptide or a fragment of at least 6 amino acids or a sequence in
XX which at least 95% of deviations from GRBP2 sequences are conservative
XX substitutions. Also included are an isolated nucleic acid (GRBP2 NA),
XX encoding GRBP2 comprising the full length cDNA or CDS, fragments or
XX variants, GRBP2 vectors, host cells, antibodies, transgenic non-human
XX animals modified to contain GRBP2 NA for unable to express the endogenous
XX orthologue of GRBP2), diagnosing a disease caused by a mutation in human
XX GRBP2 or altered expression of GRBP2, anti-agonists of GRBP2, GRBP2
XX microarrays, fusion proteins and screening for agents that modulate the
XX expression of GRBP2 NA. GRBP2 is useful for identifying binding partners
XX of GRBP2. GRBP2, GRBP2 NA and Ab are useful in therapy and in the
XX manufacture of a medicament for the treatment or prevention of a disorder
XX associated with increased or decreased expression or activity of human
XX GRBP2 (e.g. tumours, liposarcoma, ichthyosis congenita III and benign
XX familial infantile convulsion, all associated with the chromosomal
XX location of GRBP2, 19q12). GRBP2 is useful as a standard in immunoassay
XX specific for the proteins, to be used in a therapeutic agent, as
XX vaccines, to be and as antigens (e.g. for epitope mapping) or immunogens
XX (e.g. for raising antibodies). GRBP2 NA is useful as hybridisation probes,
XX to prime synthesis of nucleic acids, to prime first strand cDNA sequence
XX on an mRNA template, and to drive in vivo expression of the proteins. The
XX vector is useful for shutting GRBP2 NA between host cells derived from
XX disparate organisms, for inserting GRBP2 NA into host cell chromosome,
XX for expressing sense or antisense RNA transcripts of GRBP2 NA in vitro or
XX within a host cell, and for expressing GRBP2 alone or as fusions to
XX heterologous polypeptides. The antibody is useful as an analytical
XX reagent for detection and quantification of GRBP2 and as an immuno
XX therapeutic agent and is useful for flow cytometric detection, for
XX scanning laser cytometric detection, or for fluorescent immunoassay.
XX The present sequence is a single exon probe for GRBP2.
XX
XX Sequence 500 BP; 55 A; 163 C; 228 G; 54 T; 0 other;
XX
XX

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Query Match 10.1%; Score 50.8; DB 24; Length 500;
Best Local Similarity 45.5%; Pred. No. 0.2;
Matches 181; Conservative 0; Mismatches 217; Indels 0; Gaps 0;

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GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.
OM nucleic - nucleic search, using sw model
Run on: October 24, 2003, 13:34:49 : Search time 31.3542 Seconds
(without alignment)
7052.739 Million cell updates/sec
Title: US-09-830-902-1_COPY_10000_10500
Perfect score: 501
Sequence: 1 ggttccgcggttcgagg.....gggtcgggagggggggc ttt
Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0
Searched: 569978 seqs, 220691566 residues
Total number of hits satisfying chosen parameters: 113956

Minimum DB seq length: 0
Maximum DB seq length: 200000000
Post-processing: Minimum Match %
Listing first 45 summaries
Database : Issued Patents NAs
1: /cgn2_6/prodata/2/na/5A.CONS.seq:
2: /cgn2_6/prodata/2/na/5B.CONS.seq:
3: /cgn2_6/prodata/2/na/5C.CONS.seq:
4: /cgn2_6/prodata/2/na/6P.CONS.seq:
5: /cgn2_6/prodata/2/na/6SUS.CONS.seq:
6: /cgn2_6/prodata/2/na/backfiles1.seq:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES				Description	
Result No.	Score	Query Match	Length DB ID		
1	56.6	100	7218	US-08-232-463-14	Sequence 14, Appl
2	49.4	9.9	1269	US-08-396-218-1	Sequence 1, Appl
3	49.4	9.9	1269	US-08-760-116-1	Sequence 1, Appl
4	47.8	9.5	1735	US-08-102-863-10	Sequence 10, Appl
5	47.8	9.5	1735	PCT-US92-10895-10	Sequence 10, Appl
6	46.4	9.3	12001	US-08-458-568A-11	Sequence 11, Appl
7	46.2	9.2	2881	US-09-096-982-7	Sequence 7, Appl
8	46.2	9.2	2081	US-08-653-650A-7	Sequence 7, Appl
9	46.2	9.2	3013	US-09-096-982-6	Sequence 6, Appl
10	46.2	9.2	3013	US-08-653-650A-6	Sequence 6, Appl
11	46.2	9.2	3196	US-09-096-982-4	Sequence 4, Appl
12	46.2	9.2	3196	US-08-653-650A-4	Sequence 4, Appl
13	46	9.2	803	US-07-928-611-12	Sequence 12, Appl
14	46	9.2	803	US-08-487-811A-12	Sequence 12, Appl
15	46	9.2	803	US-09-060-694-12	Sequence 12, Appl
16	46	9.2	803	US-09-378-074-12	Sequence 12, Appl
17	46	9.2	803	PCT-US93-07370-12	Sequence 12, Appl
18	46	9.2	1504	US-09-016-434-1276	Sequence 1276, AP
19	46	9.2	1610	US-08-056-051-5	Sequence 5, Appl
20	46	9.2	1610	US-07-928-611-21	Sequence 21, Appl
21	46	9.2	1610	US-08-487-811A-21	Sequence 21, Appl
22	46	9.2	1610	US-09-060-694-21	Sequence 21, Appl
23	46	9.2	1610	US-09-378-074-21	Sequence 21, Appl
24	46	9.2	1610	PCT-US93-07370-21	Sequence 21, Appl
25	45.2	9.0	1877	US-09-780-73A-10	Sequence 10, Appl
26	45	8.8	3177	US-08-042-747A-4	Sequence 4, Appl
27	44	8.8	8438	US-07-945-283-1	Sequence 1, Appl

C 28	44	8.8	4401765	3	US-09-103-84CA-2	Sequence 2, Appl
C 29	44	8.8	4411529	3	US-09-103-84CA-1	Sequence 1, Appl
C 30	43.2	8.6	1594	4	US-09-252-991A-14592	Sequence 14590, A
C 31	43.2	8.6	1995	4	US-09-252-991A-15174	Sequence 15174, A
C 32	43.2	8.6	2109	4	US-09-252-991A-15038	Sequence 15038, A
C 33	43	8.5	2064	1	US-08-343-428-1	Sequence 1, Appl
C 34	42.6	8.5	2150	2	US-08-318-837-1	Sequence 1, Appl
C 35	42.2	8.4	4257	3	US-08-690-473-1	Sequence 1, Appl
C 36	42.2	8.4	4257	3	US-09-259-821A-1	Sequence 1, Appl
C 37	42.2	8.4	4257	3	US-08-841-659-1	Sequence 1, Appl
C 38	42.2	8.4	12001	1	US-08-458-568A-11	Sequence 11, Appl
C 39	42	8.4	1413	4	US-08-984-709A-52	Sequence 52, Appl
C 40	42	8.4	11219	4	US-07-642-734C-1	Sequence 1, Appl
C 41	42	8.4	11219	3	US-08-439-039A-1	Sequence 1, Appl
C 42	41.8	8.3	34094	4	US-09-292-034-1	Sequence 1, Appl
C 43	41.8	8.3	15233	3	US-09-128-155-16	Sequence 16, Appl
C 44	41.6	8.3	44	4	US-09-252-991A-3398	Sequence 3398, AP
C 45	41.6	5.3	2061	4	US-09-252-991A-3398	Sequence 3398, AP

ALIGNMENTS

RESULT 1
US-08-232-463-14
Sequence 14, Application US/08232463
Patent No. 5670367
GENERAL INFORMATION:
APPLICANT: CORNER, F.
APPLICANT: SCHEFFLINGER, F.
APPLICANT: FALKNER, F. G.
TITLE OF INVENTION: RECOMBINANT FOULPOX VIRUS
NUMBER OF SEQUENCES: 52
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 1800 Diagonal Road, Suite 500
City: Alexandria
STATE: VA
COUNTRY: USA
ZIP: 22333-0299
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.05
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/232,463
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/07/935,313
FILING DATE:
APPLICATION NUMBER: EP 91 114 300.6
FILING DATE: 26-AUG-1991
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 30472/114 INVU
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)836-9300
TELEFAX: (703)683-4105
TELEX: 899149
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 7218 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
CLONE: pTZgpt-Pis
US-08-232-463-14
Query Match 11.3%, Score 56.6; DB 1; Length 7218;

FILING DATE: 27-FEB-1995
ATTORNEY/AGENT INFORMATION:
NAME: KITTIS, Morice C
REGISTRATION NUMBER: 36,105
REFERENCE/DOCKET NUMBER: P1615-6007
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202/638-5000
TELEFAX: 202/638-4810
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1269 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Streptomyces peucetius
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1269
US-08-760-116-1

Query Match 9.9%; Score 49.4; DB 1; Length 1269;

Best Local Similarity 48.4%; Pred. No. 0.03;
Matches 137; Conservative 0; Mismatches 146; Indels 0; Gaps 0;

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QY 108 CAACCGGTCCTCCAGAGCTCCGCGCCCTGCTGGCGCCCGCGCCCTCCGCGCGG 167
DB 360 CGAGCGGACGATCGATCGCGGATCGCGCGGCTGTCACGAACGCGCGAGC 419
QY 168 GCGGCGCCCTCCCGGAGTGGCGGATAGCGGAACTGTACTATTTCCTCATCGCT 227
DB 420 CTCGCGCGGTCGGGCAACCGGCGAGTGAACGCGGCTTCGATACACTTCCGCT 409
QY 228 GTTTGAGATTCGCGCTGTGGCTTGTGCTGCTTCCAGCTGGGCTCCTTTGGTGG 267
DB 480 GTTGTGATTCGAGCTGTGCTGTGCGGTACCGATCGGAGATGGCGCGGAGGC 533
QY 288 GCTTCGCAAGCTTTTCGCGCGCTCATAGGAGGAAAGAGAGATCGCGAGCGGCTG 347
DB 540 CGTCAGAGTCTCAAGGACTCGGCTCGCGCGCGCGCGCGAGAGTGAAGGAC 539
QY 348 AGACATGCTCGGCTCGACCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 362
DB 600 GGACCTTCGCGGCGGTGCGGACACTTCGCGCGCTCGAGAGC 442
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RESULT 4

US-08-102-863-10

Sequence 10, Application US/08/02863

Patent No. 5468590

GENERAL INFORMATION:

APPLICANT: SARIASLANI, SIMA

TITLE OF INVENTION: CONSTITUTIVE

TITLE OF INVENTION: EXPRESSION OF P45CSOY

TITLE OF INVENTION: AND FERREDOXIN-SOY IN

TITLE OF INVENTION: STREPTOMYCES

NUMBER OF SEQUENCES: 12

CORRESPONDENCE ADDRESS:

ADDRESSEE: E. I. DU PONT DE NEMOURS

ADDRESSEE: AND COMPANY

STREET: 1007 MARKET STREET

CITY: WILMINGTON

STATE: DELAWARE

COUNTRY: USA

ZIP: 19898

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0,

SOFTWARE: Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/06/122,863
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/07/807,001
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: GALLEOS, R. THOMAS
REGISTRATION NUMBER: 32,692
REFERENCE/DOCKET NUMBER: CR-9000
TELECOMMUNICATION INFORMATION:
TELEPHONE: 302-892-7342
TELEFAX: 302-892-7949
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1735 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-06-102-863-10

Query Match 9.5%; Score 47.8; DB 1; Length 1735;

Best Local Similarity 47.3%; Pred. No. 0.068;
Matches 149; Conservative 0; Mismatches 167; Indels 0; Gaps 0;

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QY 127 CTTCCGCCCCCTTGCTGCGCCCGCCCGCTCCCGCGCGCGCGCGCGCGCG 186
DB 205 CGGCGCGCGCGAGAACCTCGACCCACCTCCCGCGCGCGCGCGCGCGCG 264
QY 197 TCCGCGCATAGCGGAACTGTACTATTTCCTACCGCTGTTTGTAGGCTTCGCG 246
DB 265 CGCGGTCGCGCTACACCGCCCGCGCGGTAGGACCGCTGCGCGAGCGCGCG 324
QY 247 CTCGCTTGGTGGCTTTCACCTGGGCTCTCTTCGCTGGCTTGGCAGGCTTC 306
DB 325 AGCGGCTACGCTCTTCGCGGACCGCCGCTCTGCGCGGTACCGGCGATGCC 384
QY 309 CGCGGCTCATGCGAGCGGAGAGAGAGTCCGCGCGCGCGCGCGCGCGCG 366
DB 385 CGTCGCTATTGCGGAGACCGCGGCTCTCCACCGACCGCGAGTTCGCGCT 444
QY 367 GCGCGCGCGCGGTCGCGCGCGCGCGCGGAGCGGAGCGGTGCGGAGTCTTC 426
DB 445 CGGCGCGAGGTTCCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 504
QY 427 TTCGAGTACATCTCC 442
DB 505 CCGGAGCACACACACC 519
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RESULT 5

PC-US92-10885-10

Sequence 10, Application PC/US9210885

GENERAL INFORMATION:

APPLICANT: SARIASLANI, SIMA

TITLE OF INVENTION: CONSTITUTIVE

TITLE OF INVENTION: EXPRESSION OF P45CSOY

TITLE OF INVENTION: AND FERREDOXIN-SOY IN

TITLE OF INVENTION: STREPTOMYCES

NUMBER OF SEQUENCES: 11

CORRESPONDENCE ADDRESS:

ADDRESSEE: E. I. DU PONT DE NEMOURS

ADDRESSEE: AND COMPANY

STREET: 1007 MARKET STREET

CITY: WILMINGTON

STATE: DELAWARE

COUNTRY: USA

ZIP: 19898

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette, 3.50 inch,

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; MEDIUM TYPE: 1.0 MB
; COMPUTER: Macintosh
; OPERATING SYSTEM: Macintosh System, 5.0
; SOFTWARE: Microsoft Word, 4.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/10985
; FILING DATE: 19921216
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: GALLEGOS, R. THOMAS
; REGISTRATION NUMBER: 32,692
; REFERENCE/DOCKET NUMBER: CR-9000-A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 302-892-7342
; TELEFAX: 302-892-7949
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1735 base pairs
; TYPE: NUCLEIC ACID
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; PCT-US92-10985-10

Query Match 9.3%; Score 47.8; DB 5; Length 1735;
Best Local Similarity 47.0%; Pred. No. 0.058;
Matches 148; Conservative 0; Mismatches 167; Indels 0; Gaps 0;

QY 127 CTTCCGCCCTTGGCTGGCCGCCGCCCTCCCGCTGGCGGCGCGCGCGCTGGCGCGGAG 186
D 127 CTTCCGCCCTTGGCTGGCCGCCGCCCTCCCGCTGGCGGCGCGCGCGCTGGCGCGGAG 186
D 205 CCGGCCCGCAGAACCTGACACCCACCTCCCGCGCGCGCGCGCGCTCTCCCGCAGGAC 264
QY 187 TCGCCGCAATAGCGGAACCTGACTATTTCTCTACCCGCTGTTTGTAGGCTTCGGGTG 246
D 187 TCGCCGCAATAGCGGAACCTGACTATTTCTCTACCCGCTGTTTGTAGGCTTCGGGTG 246
D 265 GCGGGTGGCCCTACACCGCGCGCGCGCTACGACCCCTCGCGAGCGCGCGCGGTG 324
QY 247 CTCGGTTGGTGGCTTCCACCTGGGCTCTCTCTGCTGGCTCTGGCAGCGCTTCC 306
D 247 CTCGGTTGGTGGCTTCCACCTGGGCTCTCTCTGCTGGCTCTGGCAGCGCTTCC 306
D 325 AGCGGGTCACTCTCTGACGAGCGCGCGCTGGCGGTACCGGGACGCGCTTGGCC 384
QY 307 GCGGCCCTATGACAGCGAGAGAGCTCGGCGCGCGCGCGCGAGCACTCTCGCGCTGG 366
D 307 GCGGCCCTATGACAGCGAGAGAGCTCGGCGCGCGCGCGCGAGCACTCTCGCGCTGG 366
D 385 GTCGCGTACTGCGGACCGCGCGCTCTCACGACCGCGAGCGCGGACCTTCCCGCTC 444
QY 367 GCGCGCGCGCGGTGCGGCGCGCGCGCGCGAGCGCGGTCCGAGCTTCCAGAGCGCC 426
D 367 GCGCGCGCGCGGTGCGGCGCGCGCGCGCGAGCGCGGTCCGAGCTTCCAGAGCGCC 426
D 445 GCGCGCGAGGGTTCCGCGCGCGCGCGAGCGCGCGCGCGCTCTCTCTGCGCGTCCAGC 504
QY 427 TTCGAGTACATCTCC 441
D 427 TTCGAGTACATCTCC 441
D 505 CCGGAGCAGACACC 519

RESULT 6
US-09-458-568A-11
; Sequence 11, Application US/08458568A
; Patent No. 582339
; GENERAL INFORMATION:
; APPLICANT: Schaffer, Priscilla A.
; APPLICANT: Yeh, Lily
; TITLE OF INVENTION: Compositions and Methods for Treatment of Herpesvirus
; TITLE OF INVENTION: Infections
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Woodcock, Washburn, Kurtz, Mackiewicz & No. 582339
; STREET: One Liberty Place, 46th Floor
; CITY: Philadelphia
; STATE: PA
; COUNTRY: USA
; ZIP: 19103
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible

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; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: WordPerfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/458,568A
; FILING DATE: 02-JUNE-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 06/065,146
; FILING DATE: 05-MAY-1993
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Leary Ph.D., Kathryn R.
; REGISTRATION NUMBER: 36,317
; REFERENCE/DOCKET NUMBER: DFCI-0-29
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-3100
; TELEFAX: (215) 568-3439
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1200 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
; CRIGINAL SOURCE:
; ORGANISM: Herpes simplex virus
; STRAIN: Herpes Simplex Virus Type 1
; US-38-458-568A-11

Query Match 9.3%; Score 46.4; DB 1; Length 1200;
Best Local Similarity 51.3%; Pred. No. 0.16;
Matches 158; Conservative 0; Mismatches 146; Indels 4; Gaps 2;

QY 111 CCGCGTCCCTCCGAGGCTCCGCGCGCTTGCCTGGCGCGCGCGCGCGCGCGCGCGCG 170
D 111 CCGCGTCCCTCCGAGGCTCCGCGCGCTTGCCTGGCGCGCGCGCGCGCGCGCGCGCG 170
D 1258 CCGCGTCCCTCCGCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCG 1317
QY 171 GCGCGCTCCCGCGAGTCCGCGCATAGCGGAACCTGTACTATTTTCCTACCGCTGT 230
D 171 GCGCGCTCCCGCGAGTCCGCGCATAGCGGAACCTGTACTATTTTCCTACCGCTGT 230
D 1318 CCGCGTCCCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCG 1375
QY 231 TGTAGGCTTGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 288
D 231 TGTAGGCTTGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 288
D 1376 CCGCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCG 1435
QY 231 CTCCGAGCGCTTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCG 358
D 231 CTCCGAGCGCTTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCG 358
D 1436 C-CCCCGCTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCGCGCTCTCCCG 1493
QY 351 ACCTGCTCGGCTCGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 410
D 351 ACCTGCTCGGCTCGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 410
D 1494 ACGCGCGCGGAGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 1552
QY 411 CTCCGACA 418
D 411 CTCCGACA 418
D 1554 CTCCGACA 1561

RESULT 7
US-09-096-982-7
; Sequence 7, Application US/09096982
; Patent No. 596299
; GENERAL INFORMATION:
; APPLICANT: Strobel, William R.
; APPLICANT: Dickens, Michael L.
; APPLICANT: Desanti, Charles L.
; TITLE OF INVENTION: METHOD OF PRODUCING DOXORUBICIN
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: CALFEE, HALTER & GRISMOLD
; STREET: 800 Superior Avenue, Suite 1400

```

CITY: Cleveland
STATE: Ohio
COUNTRY: USA
ZIP: 44114-2688
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/096,982
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Gollrick, Mary E.
REGISTRATION NUMBER: 34829
REFERENCE/DOCKET NUMBER: 22727/0013:
TELECOMMUNICATION INFORMATION:
TELEPHONE: 216-622-8458
TELEFAX: 216-241-0616
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2081 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 227..1649
US-09-096-982-7

Query Match 9.2%; Score 46.2; DB 2; Length 2081;
Best Local Similarity 47.7%; Pred. No. 0.15;
Matches 135; Conservative 0; Mismatches 149; Indels 0; Gaps 0;

QY 106 CAACCGGCTGCTCCAGGAGTCTGGGCGCTTTCCTGGGCGGCGGCTGCGGCGGCGG 167
DB 712 CGAGCGGCGGATCGGATCGGAGCTGCGGAGCGCTGCTGCAAGACTGGGATC 901
QY 168 GCGGGCGCTTCGCGGAGTGGCGGATAGCGGAACTGTACTATTCTCTACCGCT 227
DB 802 CTCGACCGCTGCGGCGGAGTGGCGGATAGCGGAGCTGCTGCGGAGCTGCGGAGCT 861
QY 228 GTTGTAGGCTTGGCGCTGCTGGCTTGGCTGGCTTGGCTGGCTTGGCTGGCTTGG 287
DB 862 GTTGTAGGCTTGGCGCTGCTGGCTTGGCTGGCTTGGCTGGCTTGGCTGGCTTGG 921
QY 288 GCTCTCCAGCGCTTTCGCGGCGGCTGATGCGAGCGGAGAGAGCTCGGCGGCGG 347
DB 922 CGTCCGCGTCTCAAGGACTTGGCGCTGCGGCGGCGGAGAGCGGCGGCGTACGCGC 981
QY 348 AGCAGCTGCTCGGCTGCTGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGG 390
DB 982 GGACCTGCGGCGGAGCTGCGGAGAGCTGCGGAGAGCTGCGGAGAGC 1024

RESULT 8
US-09-096-982-7
Sequence 7, Application US/08653650A
Patent No. 5978830
GENERAL INFORMATION:
APPLICANT: Strohl, William P.
APPLICANT: Dickens, Michael L.
APPLICANT: Desanti, Charles L.
TITLE OF INVENTION: METHOD OF PRODUCING DOXORUBICIN
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: CALFEE, HALTER & GRISWOLD
STREET: 800 Superior Avenue, Suite 1400
CITY: Cleveland
STATE: Ohio
COUNTRY: USA

ZIP: 44114-2688
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/653,650A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Gollrick, Mary E.
REGISTRATION NUMBER: 34829
REFERENCE/DOCKET NUMBER: 22727/0013:
TELECOMMUNICATION INFORMATION:
TELEPHONE: 216-622-8458
TELEFAX: 216-241-0616
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2081 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 227..1649
US-09-653-650A-7

Query Match 9.2%; Score 46.2; DB 2; Length 2081;
Best Local Similarity 47.7%; Pred. No. 0.15;
Matches 135; Conservative 0; Mismatches 149; Indels 0; Gaps 0;

QY 108 CAACCGGCTGCTCCAGGAGTCTGGGCGCTTTCCTGGGCGGCGGCTGCGGCGGCGG 167
DB 742 CGAGCGGCGGATCGGATCGGAGCTGCGGAGCGCTGCTGCAAGACTGGGATC 801
QY 168 GCGGGCGCTTCGCGGAGTGGCGGATAGCGGAACTGTACTATTCTCTACCGCT 227
DB 802 CTCGACCGCTGCGGCGGAGTGGCGGATAGCGGAGCTGCTGCGGAGCTGCGGAGCT 861
QY 228 GTTGTAGGCTTGGCGCTGCTGGCTTGGCTGGCTTGGCTGGCTTGGCTGGCTTGG 287
DB 862 GTTGTAGGCTTGGCGCTGCTGGCTTGGCTGGCTTGGCTGGCTTGGCTGGCTTGG 921
QY 288 GCTCTCCAGCGCTTTCGCGGCGGCTGATGCGAGCGGAGAGAGCTCGGCGGCGG 347
DB 922 CGTCCGCGTCTCAAGGACTTGGCGCTGCGGCGGCGGAGAGCGGCGGCGTACGCGC 981
QY 348 AGCAGCTGCTCGGCTGCTGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGG 390
DB 982 GGACCTGCGGCGGAGCTGCGGAGAGCTGCGGAGAGCTGCGGAGAGC 1024

RESULT 9
US-09-096-982-6
Sequence 6, Application US/09096982
Patent No. 5962293
GENERAL INFORMATION:
APPLICANT: Strohl, William R.
APPLICANT: Dickens, Michael L.
APPLICANT: Desanti, Charles L.
TITLE OF INVENTION: METHOD OF PRODUCING DOXORUBICIN
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: CALFEE, HALTER & GRISWOLD
STREET: 800 Superior Avenue, Suite 1400
CITY: Cleveland
STATE: Ohio
COUNTRY: USA
ZIP: 44114-2688
COMPUTER READABLE FORM: disk
MEDIUM TYPE: Floppy

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COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/096,982
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Goirick, Mary E.
REGISTRATION NUMBER: 34829
REFERENCE/DOCKET NUMBER: 22727/00131
TELECOMMUNICATION INFORMATION:
TELEPHONE: 216-622-8458
TELEFAX: 216-241-0816
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 3013 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
US-09-096-982-6

Query Match
Best Local Similarity 47.7%; Pred. No. 0.16;
Matches 135; Conservative 0; Mismatches 144; Indels 0; Gaps 0;

CY 108 CAACCGGGTGGCTTCCGAGGAGTCCGCGGCTTACCTGGAGGCGGCGGCGGCGGCGG 167
DB 1674 CGAGCGGAGCGATCGCATCGCGGCGATCGCGGCGATCGCGGCGATCGCGGCGATC 1733
CY 168 GCGGGCGGCTTCCGCGGAGTCCGCGGATAGCGGAGACCTGTACTATTTCCTACCGGCT 227
DB 1734 CTCGACCGGTTCGGGGAACCGCGGAGCTGATCGCGGCTTCCGCTACCACTTCCGCT 1793
CY 228 GTTTGTAGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCT 287
DB 1794 GTTGTGATCTCGAATCTCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCG 1853
CY 288 GCTTCCGAGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCT 347
DB 1854 CGTGGCGGTGCTCAAGGCACTCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCT 1913
CY 348 AGCACTGCTCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCT 390
DB 1914 GGACCTTCCGCGGAGCTGCGCGGAGACGTCGCGGCTTCCGCGGCTTCCGCGGCT 1956
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RESULT 10
US-08-653-650A-6
Sequence 6, Application US/08653650A
Patent No. 5976830
GENERAL INFORMATION:
APPLICANT: Strohl, William R.
APPLICANT: Dickens, Michael L.
APPLICANT: Desanti, Charles L.
TITLE OF INVENTION: METHOD OF PRODUCING DOXORUBICIN
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: CALFE, HAUTER & GRISWOLD
STREET: 800 Superior Avenue, Suite 1400
CITY: Cleveland
STATE: Ohio
COUNTRY: USA
ZIP: 44114-2689
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/653,650A
FILING DATE:
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CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Goirick, Mary E.
REGISTRATION NUMBER: 34829
REFERENCE/DOCKET NUMBER: 22727/00131
TELECOMMUNICATION INFORMATION:
TELEPHONE: 216-622-8458
TELEFAX: 216-241-0816
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 3013 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
US-08-653-650A-6

Query Match
Best Local Similarity 47.7%; Pred. No. 0.16;
Matches 135; Conservative 0; Mismatches 148; Indels 0; Gaps 0;

CY 108 CAACCGGGTGGCTTCCGAGGAGTCCGCGGCTTACCTGGAGGCGGCGGCGGCGGCGG 167
DB 1674 CGAGCGGAGCGATCGCATCGCGGCGATCGCGGCGATCGCGGCGATCGCGGCGATC 1733
CY 168 GCGGGCGGCTTCCGCGGAGTCCGCGGATAGCGGAGACCTGTACTATTTCCTACCGGCT 227
DB 1734 CTCGACCGGTTCGGGGAACCGCGGAGCTGATCGCGGCTTCCGCTACCACTTCCGCT 1793
CY 228 GTTTGTAGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCT 287
DB 1794 GTTGTGATCTCGAATCTCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCG 1853
CY 288 GCTTCCGAGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCT 347
DB 1854 CGTGGCGGTGCTCAAGGCACTCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCT 1913
CY 348 AGCACTGCTCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCTTCCGCGGCT 390
DB 1914 GGACCTTCCGCGGAGCTGCGCGGAGACGTCGCGGCTTCCGCGGCTTCCGCGGCT 1956

RESULT 11
US-09-096-982-4
Sequence 4, Application US/09096982
Patent No. 5962293
GENERAL INFORMATION:
APPLICANT: Strohl, William R.
APPLICANT: Dickens, Michael L.
APPLICANT: Desanti, Charles L.
TITLE OF INVENTION: METHOD OF PRODUCING DOXORUBICIN
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: CALFE, HAUTER & GRISWOLD
STREET: 800 Superior Avenue, Suite 1400
CITY: Cleveland
STATE: Ohio
COUNTRY: USA
ZIP: 44114-2689
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/096,982
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Goirick, Mary E.
REGISTRATION NUMBER: 34829
REFERENCE/DOCKET NUMBER: 22727/00131
TELECOMMUNICATION INFORMATION:
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TELEPHONE: 216-622-8458
TELEFAX: 216-241-0816
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 3196 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: mat peptide
LOCATION: 1495..2764
FEATURE:
NAME/KEY: CDS
LOCATION: 1495..2764
US-09-096-982-4

Query Match 9.2% Score 46.2; DB 2; Length 3196;
Best Local Similarity 47.7% Pred. No. 0.16;
Matches 135; Conservative 0; Mismatches 148; Indels 0; Gaps 0;

QY 108 CAACCGGCTGCTCCAGAGCTCCGCGCCCTTGGCTGAGCCCGGCGCTCCGCGCGCG 167
DB 1857 CGAGCGAGGATCGCATCGCGCCCATCGCGACCGCTGCTCAGCAACTCGCGACTC 1916
QY 168 GCGGCGCCCTCCGCGAGTCGCGCGATAGCGGACCTGTACTATTCTCTACCGCT 227
DB 1917 CTCGACCGGCTCGGCGAACCGCGCGAGCTGATCGGCGCTTCGGGTACCACTTCGCGT 1976
QY 228 GTTTGAGGCTTCGCGCTGCTGGCTTGGCTTGGCTTCCAGCTGGGCTCTCTTCTGTG 287
DB 1977 GTTGGTCATCTCGCACTCTCGGCTGCGGTCGCGTCTGCGGATCGCGCGGAGGC 2018
QY 288 GCTCTGCAAGCGCTTTCGCGCGCTCTATGCGAGCGACAGAGAGCTCGGCGCGCGCG 347
DB 2037 CGTGCAGCGCTCAAGGCACTCGGCTTGGCGCGCGAGAGCGCGGCTGACGCGAC 2096
QY 348 AGACCTGCTCGCGCTTCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 390
DB 2097 GACCTTCGCGGAGCTGTCGCGACAGCTCGCGCTGGAGAGC 2139

RESULT 12
US-08-653-650A-4
Sequence 4, Application US/08653650A
Patent No. 5976830
GENERAL INFORMATION:
APPLICANT: Strohl, William P.
APPLICANT: Dickens, Michael L.
APPLICANT: Desanti, Charles L.
TITLE OF INVENTION: METHOD OF PRODUCING DOXORUBICIN
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: CALPEE, HAUTER & GRISWOLD
STREET: 800 Superior Avenue, Suite 1400
City: Cleveland
STATE: Ohio
COUNTRY: USA
ZIP: 44114-2688
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/653,650A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Geilick, Mary E.
REGISTRATION NUMBER: 14829
REFERENCE/DOCKET NUMBER: 22727/00131
TELECOMMUNICATION INFORMATION:

TELEPHONE: 216-622-8458
TELEFAX: 216-241-0816
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 3196 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: mat peptide
LOCATION: 1495..2764
FEATURE:
NAME/KEY: CDS
LOCATION: 1495..2764
US-08-653-650A-4

Query Match 9.2% Score 46.2; DB 2; Length 3196;
Best Local Similarity 47.7% Pred. No. 0.16;
Matches 135; Conservative 0; Mismatches 148; Indels 0; Gaps 0;

QY 108 CAACCGGCTGCTCCAGAGCTCCGCGCCCTTGGCTGAGCCCGGCGCTCCGCGCGCG 167
DB 1857 CGAGCGAGGATCGCATCGCGCCCATCGCGACCGCTGCTCAGCAACTCGCGACTC 1916
QY 168 GCGGCGCCCTCCGCGAGTCGCGCGATAGCGGACCTGTACTATTCTCTACCGCT 227
DB 1917 CTCGACCGGCTCGGCGAACCGCGCGAGCTGATCGGCGCTTCGGGTACCACTTCGCGT 1976
QY 228 GTTTGAGGCTTCGCGCTGCTGGCTTGGCTTGGCTTCCAGCTGGGCTCTCTTCTGTG 287
DB 1977 GTTGGTCATCTCGCACTCTCGGCTGCGGTCGCGTCTGCGGATCGCGCGGAGGC 2018
QY 288 GCTCTGCAAGCGCTTTCGCGCGCTCTATGCGAGCGACAGAGAGCTCGGCGCGCGCG 347
DB 2037 CGTGCAGCGCTCAAGGCACTCGGCTTGGCGCGCGAGAGCGCGGCTGACGCGAC 2096
QY 348 AGACCTGCTCGCGCTTCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 390
DB 2097 GACCTTCGCGGAGCTGTCGCGACAGCTCGCGCTGGAGAGC 2139

RESULT 13
US-07-928-611-12
Sequence 12, Application US/07928611
Patent No. 5565601
GENERAL INFORMATION:
APPLICANT: Van To, Hubert H.M.
APPLICANT: Civeili, Olivier
TITLE OF INVENTION: A No. 5565601a: Human Dopamine Receptor and Uses
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Allegretti & Witcoff, Ltd.
STREET: 10 South Wacker Drive, Suite 3000
City: Chicago
STATE: Illinois
COUNTRY: USA
ZIP: 60606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/928,611
FILING DATE: 19920810
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: No. 5569601nan, Kevin E
REGISTRATION NUMBER: 35,303
REFERENCE/DOCKET NUMBER: 90,1092-B
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312-715-1000

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Run on: October 24, 2003, 15:28:54 ; Search time 123.415 Seconds
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Title: US-09-830-902-1_COPY_10000_10500

Perfect score: 501
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Scoring table: IDENTITY NJC
Gapop 10.0, Gapext 1.0

Searched: 1792395 seqs, 1345950451 residues

Total number of hits satisfying chosen parameters: 3584730

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:

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- 2: /cgn2_6/prodata/2/pubpra/PCT_NEW_PUB.seq.
- 3: /cgn2_6/prodata/2/pubpra/US06_NEW_PUB.seq.
- 4: /cgn2_6/prodata/2/pubpra/US06_PUBCOMB.seq.
- 5: /cgn2_6/prodata/2/pubpra/US07_NEW_PUB.seq.
- 6: /cgn2_6/prodata/2/pubpra/PCTUS_PUBCOMB.seq.
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- 8: /cgn2_6/prodata/2/pubpra/US06_PUBCOMB.seq.
- 9: /cgn2_6/prodata/2/pubpra/US05A_PUBCOMB.seq.
- 10: /cgn2_6/prodata/2/pubpra/US06_PUBCOMB.seq.
- 11: /cgn2_6/prodata/2/pubpra/US09_PUBCOMB.seq.
- 12: /cgn2_6/prodata/2/pubpra/US09_NEW_PUB.seq.
- 13: /cgn2_6/prodata/2/pubpra/US10A_PUBCOMB.seq.
- 14: /cgn2_6/prodata/2/pubpra/US10B_PUBCOMB.seq.
- 15: /cgn2_6/prodata/2/pubpra/US10_NEW_PUB.seq.
- 16: /cgn2_6/prodata/2/pubpra/US06_NEW_PUB.seq.
- 17: /cgn2_6/prodata/2/pubpra/US06_PUBCOMB.seq.

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	48.2	9.5	956	13	US-10-027-632-31508
3	47.4	9.5	1117	12	US-10-017-161-1403
C 4	47	9.4	594	12	US-10-140-472-10
C 5	47	9.4	594	12	US-10-141-765-10
C 6	47	9.4	594	12	US-10-142-895-10
C 7	47	9.4	594	12	US-10-158-795-10
C 8	47	9.4	594	14	US-10-123-155-10
C 9	47	9.4	594	15	US-10-146-731-10
C 10	47	9.4	3133	12	US-10-017-161-1483
11	46.6	9.3	31168	10	US-09-764-868-1464
12	46.4	9.3	1719	14	US-10-156-761-5084
13	46.4	9.3	9025608	14	US-10-156-761-1
14	46	9.2	803	14	US-10-224-265-12
15	46	9.2	1504	14	US-10-225-567A-105
16	46	9.2	1610	14	US-10-224-265-21

C 17	45.2	9.0	2364	14	US-10-156-761-3756	Sequence 3756, Ap
C 18	44.6	8.9	1374	14	US-10-156-761-5859	Sequence 5859, Ap
C 19	44.4	8.9	3163	12	US-10-017-161-1857	Sequence 1857, Ap
C 20	44.4	8.9	5452	12	US-10-017-161-1481	Sequence 1481, Ap
C 21	44.2	8.8	154746	12	US-09-827-688-8	Sequence 8, Appl
C 22	44.2	8.8	154746	12	US-09-827-688-8	Sequence 8, Appl
C 23	43.8	8.7	732	12	US-10-259-165-115	Sequence 115, App
C 24	43.6	8.7	3059	12	US-10-007-826A-408	Sequence 408, App
C 25	43.4	8.7	12350	14	US-10-156-761-3627	Sequence 3627, Ap
C 26	43.4	8.7	1614	10	US-03-976-743-45	Sequence 45, App
C 27	43.4	8.7	1614	13	US-10-023-529-45	Sequence 45, App
C 28	43.4	8.7	1514	13	US-10-023-523-45	Sequence 45, App
C 29	43.4	8.7	12425	13	US-03-976-740-50	Sequence 50, Appl
C 30	43.4	8.7	12425	13	US-10-023-529-50	Sequence 50, Appl
C 31	43.4	8.7	12425	13	US-10-023-523-50	Sequence 50, Appl
C 32	43.4	8.7	89421	10	US-09-976-059-1	Sequence 1, Appl
C 33	43.4	8.7	9025608	14	US-10-156-761-1	Sequence 1, Appl
C 34	43	8.6	2307	11	US-09-933-512A-87	Sequence 87, Appl
C 35	43	8.6	3670	14	US-10-074-475-98	Sequence 98, Appl
C 36	42.8	8.5	1951	10	US-09-736-368A-104	Sequence 104, App
C 37	42.8	8.5	7771	10	US-09-832-352-38	Sequence 38, Appl
C 38	42.6	8.5	1254	12	US-10-393-593-38	Sequence 38, Appl
C 39	42.6	8.5	1254	12	US-10-393-567-38	Sequence 38, Appl
C 40	42.6	8.5	1254	12	US-10-394-087-38	Sequence 38, Appl
C 41	42.6	8.5	1258	14	US-10-171-581-199	Sequence 199, App
C 42	42.4	8.5	1199	12	US-10-029-386-25427	Sequence 25427, A
C 43	42.4	8.5	1866	11	US-09-945-327-51	Sequence 51, Appl
C 44	42.4	8.5	2124	10	US-09-815-925-5	Sequence 8, Appl
C 45	42.2	8.4	1786	13	US-10-027-632-37688	Sequence 97688, A

ALIGNMENTS

RESULT 1
US-09-895-040A-23
; Sequence 23, Application US/09895040A
; Patent No. US20020123474A1
; GENERAL INFORMATION:
; APPLICANT: Shantou, Mark
; APPLICANT: Ji, Yonggang
; TITLE OF INVENTION: HUMAN GTP-RHO BINDING PROTEIN 2
; FILE REFERENCE: AEMICA-11
; CURRENT APPLICATION NUMBER: US/09/895,040A
; PRIORITY FILING DATE: 2001-06-29
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIORITY FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIORITY FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIORITY FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIORITY FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIORITY FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIORITY FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIORITY FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIORITY FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIORITY FILING DATE: 2001-05-23
; NUMBER OF SEQ ID NOS: 180
; SOFTWARE: Aemica Sequence Listing Engine
; SEQ ID NO: 23
; LENGTH: 500
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-895-040A-23

Query Match 10.1%; Score 50.8; DB 10; Length 500;
Best Local Similarity 45.5%; Pred. No. 0.00371;

FEATURE:	NAME/KEY: modified_base	
	LOCATION: (552)	
OTHER INFORMATION:	a, t, c, g, unknown or other	
FEATURE:	NAME/KEY: modified_base	
	LOCATION: (557)..(561)	
OTHER INFORMATION:	a, t, c, g, unknown or other	
FEATURE:	NAME/KEY: modified_base	
	LOCATION: (565)	
OTHER INFORMATION:	a, t, c, g, unknown or other	
FEATURE:	NAME/KEY: modified_base	
	LOCATION: (573)	
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FEATURE:	NAME/KEY: modified_base	
	LOCATION: (577)	
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FEATURE:	NAME/KEY: modified_base	
	LOCATION: (588)	
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FEATURE:	NAME/KEY: modified_base	
	LOCATION: (605)..(610)	
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FEATURE:	NAME/KEY: modified_base	
	LOCATION: (617)	
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	LOCATION: (619)..(622)	
OTHER INFORMATION:	a, t, c, g, unknown or other	
FEATURE:	NAME/KEY: modified_base	
	LOCATION: (629)	
OTHER INFORMATION:	a, t, c, g, unknown or other	
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OTHER INFORMATION:	a, t, c, g, unknown or other	
FEATURE:	NAME/KEY: modified_base	
	LOCATION: (648)	
OTHER INFORMATION:	a, t, c, g, unknown or other	
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	LOCATION: (650)	
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	LOCATION: (652)..(653)	
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OTHER INFORMATION:	a, t, c, g, unknown or other	
FEATURE:	NAME/KEY: modified_base	
	LOCATION: (688)..(689)	
OTHER INFORMATION:	a, t, c, g, unknown or other	
FEATURE:	NAME/KEY: modified_base	


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QY 288 GCTCTGCCAGCGCTTCTCCGCGCGCTCATGCGAGGAGAGAGAGCTCCGCGCGCGCG 347
DB 390 SAYSTSSSSSSSYVYTSNYC.T.CC...T.MCAABCS...TTTTTTT...HSCC.S 331
QY 342 AGCACTGCTCCGCTCCGCGCGCGCGCGCTGCGGCGGAGAGAGAGAGAGAGAGAG 407
DB 332 A..A.M..YC.A.SYSYS.SSS.S.SYMR.HRA.SHYTRS..S.MYCY.YY.Y.YY 271
QY 408 AGCTTTCCA 416
DB 270 .YVSYYCSR 262

RESULT 6
US-10-142-985-10/C
; Sequence 10, Application US/10142985
; Publication No. US2003015764A1
; GENERAL INFORMATION:
; APPLICANT: Baker, Kevin P.
; APPLICANT: Beresini, Maureen
; APPLICANT: DeForge, Laura
; APPLICANT: Desnoyers, Luc
; APPLICANT: Filvaroff, Ellen
; APPLICANT: Gao, Wei-Qiang
; APPLICANT: Gerritsen, Mary E.
; APPLICANT: Goddard, Audrey
; APPLICANT: Godowski, Paul C.
; APPLICANT: Gurney, Austin L.
; APPLICANT: Sherwood, Steven
; APPLICANT: Smith, Victoria
; APPLICANT: Stewart, Timothy A.
; APPLICANT: Tomas, Daniel
; APPLICANT: Watanabe, Colin K
; APPLICANT: Wood, William
; APPLICANT: Zhang, Zemin
; TITLE OF INVENTION: SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC
; FILE REFERENCE: P333R1C248
; CURRENT APPLICATION NUMBER: US/10/142,855
; CURRENT FILING DATE: 2002-05-13
; Prior Application removed - See File Wrapper or Palm
; NUMBER OF SEQ ID NOS: 562
; SEQ ID NO 10
; LENGTH: 594
; TYPE: PRT
; ORGANISM: Homo Sapien
US-10-142-985-10

Query Match 9.4%; Score 47; DB 12; Length 594;
Best Local Similarity 6.8%; Pred. No. 0.0067;
Matches 21; Conservative 152; Mismatches 132; Indels 4; Gaps 1;

QY 108 CAACCCGGTCCCTCCAGAGAGTCCGCGCGCTTCCCTCCGCGCGCGCGCGCGCGCG 167
DB 566 CA.MCTT..AYM.N.CBT..STT.YA.M.YT.S.S.S.SYSYSYS.S.S.SYSYA.S 507
QY 168 GCGCGCCCTCCGCGCGCGCGCATAGCGAGAGCTGTACTATTTCTCTACCGCT 227
DB 506 YSYS.S.SWSYSYSYSSDDY.CYCCYVRYHCSDSYSYVY.C---RCYVY.SYSRYD 451
QY 228 GTTGTAGGCTTCGGGCTGCTGGCTTGGCTTCACCTGGGCTCCTCTCTGTGTG 287
DB 450 CHYSCCSDYCYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYS 391
QY 288 GTCTGCCAGCGCTTCTCCGCGCGCGCTCATGCGAGAGAGAGAGAGCTCCGCGCGCG 347
DB 390 SAYSTSSSSSSSYVYTSNYC.T.CC...T.MCAABCS...TTTTTTT...HSCC.S 331
QY 342 AGCACTGCTCCGCTCCGCGCGCGCGCTGCGGCGGAGAGAGAGAGAGAGAGAG 407
DB 332 A..A.M..YC.A.SYSYS.SSS.S.SYMR.HRA.SHYTRS..S.MYCY.YY.Y.YY 271
QY 408 AGCTTTCCA 416
DB 270 .YVSYYCSR 262

RESULT 8
US-10-123-155-10/C
; Sequence 10, Application US/10123155
; Publication No. US2003006879A1

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QY 408 AGCTTTCCA 416
DB 270 .YVSYYCSR 262

RESULT 7
US-10-158-790-10/C
; Sequence 10, Application US/10158790
; Publication No. US20030180879A1
; GENERAL INFORMATION:
; APPLICANT: Baker, Kevin P.
; APPLICANT: Beresini, Maureen
; APPLICANT: DeForge, Laura
; APPLICANT: Desnoyers, Luc
; APPLICANT: Filvaroff, Ellen
; APPLICANT: Gao, Wei-Qiang
; APPLICANT: Gerritsen, Mary E.
; APPLICANT: Goddard, Audrey
; APPLICANT: Godowski, Paul C.
; APPLICANT: Gurney, Austin L.
; APPLICANT: Sherwood, Steven
; APPLICANT: Smith, Victoria
; APPLICANT: Stewart, Timothy A.
; APPLICANT: Tomas, Daniel
; APPLICANT: Watanabe, Colin K
; APPLICANT: Wood, William
; APPLICANT: Zhang, Zemin
; TITLE OF INVENTION: SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC
; FILE REFERENCE: P333R1C448
; CURRENT APPLICATION NUMBER: US/10/158,790
; CURRENT FILING DATE: 2002-05-30
; Prior Application removed - See File Wrapper or Palm
; NUMBER OF SEQ ID NOS: 550
; SEQ ID NO 10
; LENGTH: 594
; TYPE: PRT
; ORGANISM: Homo Sapien
US-10-158-790-10

Query Match 9.4%; Score 47; DB 12; Length 594;
Best Local Similarity 6.8%; Pred. No. 0.0067;
Matches 21; Conservative 152; Mismatches 132; Indels 4; Gaps 1;

QY 108 CAACCCGGTCCCTCCAGAGTCCGCGCGCTTCCCTCCGCGCGCGCGCGCGCGCG 167
DB 566 CA.MCTT..AYM.N.CBT..STT.YA.M.YT.S.S.S.SYSYSYS.S.S.SYSYA.S 507
QY 168 GCGCGCCCTCCGCGCGCGCGCATAGCGAGAGCTGTACTATTTCTCTACCGCT 227
DB 506 YSYS.S.SWSYSYSYSSDDY.CYCCYVRYHCSDSYSYVY.C---RCYVY.SYSRYD 451
QY 228 GTTGTAGGCTTCGGGCTGCTGGCTTGGCTTCACCTGGGCTCCTCTCTGTGTG 287
DB 450 CHYSCCSDYCYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYSYS 391
QY 288 GTCTGCCAGCGCTTCTCCGCGCGCGCTCATGCGAGAGAGAGAGAGCTCCGCGCGCG 347
DB 390 SAYSTSSSSSSSYVYTSNYC.T.CC...T.MCAABCS...TTTTTTT...HSCC.S 331
QY 342 AGCACTGCTCCGCTCCGCGCGCGCGCTGCGGCGGAGAGAGAGAGAGAGAGAG 407
DB 332 A..A.M..YC.A.SYSYS.SSS.S.SYMR.HRA.SHYTRS..S.MYCY.YY.Y.YY 271
QY 408 AGCTTTCCA 416
DB 270 .YVSYYCSR 262

RESULT 8
US-10-123-155-10/C
; Sequence 10, Application US/10123155
; Publication No. US2003006879A1

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GENERAL INFORMATION:

APPLICANT: Baker, Kevin P.

APPLICANT: Beresini, Maureen

APPLICANT: DeForge, Laura

APPLICANT: Desnoyers, Luc

APPLICANT: Filvaroff, Ellen

APPLICANT: Gao, Wei-Qiang

APPLICANT: Gerritsen, Mary E.

APPLICANT: Goddard, Audrey

APPLICANT: Godowski, Paul J.

APPLICANT: Gurney, Austin L.

APPLICANT: Sherwood, Steven

APPLICANT: Smith, Victoria

APPLICANT: Stewart, Timothy A.

APPLICANT: Tumas, Daniel

APPLICANT: Watanabe, Colin K

APPLICANT: Wood, William

APPLICANT: Zhang, Zemin

TITLE OF INVENTION: SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC

FILE REFERENCE: P3330R1C33

CURRENT APPLICATION NUMBER: US/10/123,155

CURRENT FILING DATE: 2002-04-15

Prior Application removed - See Palm or File Wrapper

NUMBER OF SEQ ID NOS: 550

SEQ ID NO 10

LENGTH: 594

TYPE: PRT

ORGANISM: Homo Sapien

US-10-123-155-10

Query Match

Best Local Similarity 9.4%; Score 47; DB 14; Length 594;

Sequence 1483; Application US/10017161

Publication No. US20030143669A2

GENERAL INFORMATION:

APPLICANT: SUWA, MAKIKO

APPLICANT: ASAI, KIYOSHI

APPLICANT: ARIYAMA, YU-AKA

APPLICANT: ABURATANI, HIROYUKI

TITLE OF INVENTION: NOVEL G PROTEIN-COUPLED RECEPTORS

FILE REFERENCE: 084335/0152

CURRENT APPLICATION NUMBER: US/10/017,161

CURRENT FILING DATE: 2002-12-18

PRIOR APPLICATION NUMBER: JP 2301/246789

PRIOR FILING DATE: 2001-06-18

NUMBER OF SEQ ID NOS: 2430

SOFTWARE: PatentIn Ver. 2.1

SEQ ID NO 1483

LENGTH: 3133

TYPE: DNA

ORGANISM: Homo sapiens

FEATURE:

US-10-146-731-10/C

Sequence 10; Application US/10146731

Publication No. US20030129692A2

GENERAL INFORMATION:

APPLICANT: Baker, Kevin P.

APPLICANT: Beresini, Maureen

APPLICANT: DeForge, Laura

APPLICANT: Desnoyers, Luc

APPLICANT: Filvaroff, Ellen

APPLICANT: Gao, Wei-Qiang

APPLICANT: Gerritsen, Mary E.

APPLICANT: Goddard, Audrey

APPLICANT: Godowski, Paul J.

APPLICANT: Gurney, Austin L.

APPLICANT: Sherwood, Steven

APPLICANT: Smith, Victoria

APPLICANT: Stewart, Timothy A.

APPLICANT: Tumas, Daniel

APPLICANT: Watanabe, Colin K

APPLICANT: Wood, William

APPLICANT: Zhang, Zemin

TITLE OF INVENTION: SECRETED AND TRANSMEMBRANE POLYPEPTIDES AND NUCLEIC

FILE REFERENCE: P3330R1C33

CURRENT APPLICATION NUMBER: US/10/123,155

CURRENT FILING DATE: 2002-04-15

Prior Application removed - See Palm or File Wrapper

NUMBER OF SEQ ID NOS: 550

SEQ ID NO 10

LENGTH: 594

TYPE: PRT

ORGANISM: Homo Sapien

US-10-123-155-10

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NAME/KEY: CDS
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NAME/KEY: modified_base
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OTHER INFORMATION: a, t, c, g, unknown or other
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FEATURE:
NAME/KEY: modified_base
LOCATION: (370)
OTHER INFORMATION: a, t, c, g, unknown or other
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RESULT 13

US-10-156-761-1

Sequence 1, Application: US/015576;

Publication No. US2003019018A1

GENERAL INFORMATION:

APPLICANT: OMURA, SATOSHI

APPLICANT: KEDA, HARUO

APPLICANT: ISHIKAWA, JUN

APPLICANT: HORIKAWA, HIROSHI

APPLICANT: SHIBA, TADAYOSHI

APPLICANT: SAKAKI, YOSHIYUKI

APPLICANT: HATTORI, YASAHISA

TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES

FILE REFERENCE: 349-262

CURRENT APPLICATION NUMBER: US/20/155,761

CURRENT FILING DATE: 2002-05-29

PRIOR APPLICATION NUMBER: JP 2001-204289

PRIOR FILING DATE: 2001-05-30

PRIOR APPLICATION NUMBER: JP 2001-272697

PRIOR FILING DATE: 2001-08-02

NUMBER OF SEQ ID NOS: 15:09

SEQ ID NO 1

LENGTH: 9025608

TYPE: DNA

ORGANISM: Streptomyces avermitilis

FEATURE:

NAME/KEY: misc feature

LOCATION: (4187715)

OTHER INFORMATION: a, t, c, g, other or unknown

US-10-156-761-1

Query Match 9.3%; Score 46.4; DB 14; Length 9025608;

Best Local Similarity 52.0%; Pred. No. 3.01;

Matches 104; Conservative 0; Mismatches 36; Indels 2; Gaps 0;

QY 257 TCSCCTTCACCTGGGGGCTCTTTCTGTGGCTGTGCAGAGCTTCTCTCCGGGCGGCA 316

DB 6193898 TCGGCGGGGCACTCCGCTGTCTGGCGGGGGCTGTGGCTGTGCAGAGCTGTGCAGGCGAG 6191957

QY 317 TGGCAGCGAGAGAGAGCTCGGGGGCGCGCCACAGCTGGCTGGGCTGGGCGGCGGCG 376

DB 6193958 TCSCCGCGGGAGGAGGCTGTGGGGGCTGTGGCCCGGATCGAGACCGCGGAGGCG 6194017

QY 377 CGGTGGCGGGCGGGAGGCGGAGCGGTTCGACATCTTCCACAAAGCGGCTTGAGTACA 416

DB 6194018 CCGCGGTGCGCGGGGGCGGCGAGCGGAGGCTGTGGCGGAGGCGGGGCGGTACG 6194077

QY 437 TCTCCATTGCGCTCGGCATC 456

DB 6194078 ACCCGAACGGCGGCACTAGC 6194087

RESULT 14

US-10-224-260-12

Sequence 12, Application: US/10224260

Publication No. US20030059845A1

GENERAL INFORMATION:

APPLICANT: Van Tol, Hubert H.M.

APPLICANT: Civelis, Olivier

TITLE OF INVENTION: A No. US20030059845A1:el Human Dopamine Receptor and Uses

NUMBER OF INVENTIONS: 22

CORRESPONDENCE ADDRESS:

ADDRESSEE: Allegretti & Witcoff, Ltd.

STREET: 10 South Wacker Drive, Suite 3000

City: Chicago

STATE: Illinois

COUNTRY: USA

ZIP: 60606

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA: US/10/224,260

APPLICATION NUMBER: US/10/224,260

FILING DATE: 20-Aug-2002

CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 09/928,611

FILING DATE: <Unknown>

ATTORNEY/AGENT INFORMATION:

NAME: No. US20030059845A1:ran, Kevin E

REGISTRATION NUMBER: 35,303

REFERENCE/DOCKET NUMBER: 90,1092-B

TELECOMMUNICATION INFORMATION:

TELEPHONE: 312-715-1000

TELEFAX: 312-715-1234

TELEX: 810-221-8117

INFORMATION FOR SEQ ID NO: 12:

SEQUENCE CHARACTERISTICS:

LENGTH: 803 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

FEATURE:

NAME/KEY: exon

LOCATION: 1..803

IDENTIFICATION METHOD: experimental

OTHER INFORMATION: /evidence= EXPERIMENTAL

/standard name: "Alternate Exon 3: D4.7"

/note: "This sequence represents the third exon of

allele D4.7 of the human D4 dopamine receptor

Gene"

FEATURE:

NAME/KEY: misc feature

LOCATION: 257..262

IDENTIFICATION METHOD: experimental

OTHER INFORMATION: /function= "PstI site"

/evidence= EXPERIMENTAL

/standard name: "PstI site"

/label= PstI

/note: "This sequence is a PstI site whereby

digestion of human genomic DNA produces a RFLP"

FEATURE:

NAME/KEY: repeat_region

LOCATION: 346..352

IDENTIFICATION METHOD: experimental

OTHER INFORMATION: /rpt_type= "tandem"

/evidence= EXPERIMENTAL

/rpt_unit= 346..354

/note: "This sequence is a repeat found in 7 known

alleles of the human D4 dopamine receptor gene

encoding a 16 amino acid sequence repeated 7 times

FEATURE:

NAME/KEY: CDS

LOCATION: 2..803

SEQUENCE DESCRIPTION: SEQ ID NO: 12:

US-10-224-260-12

Query Match

Best Local Similarity 49.7%; Pred. No. 2.012;

Matches 145; Conservative 0; Mismatches 145; Indels 2; Gaps 1;

QY 117 GGTTCACAGGCTTCGCGCGGCTTGGCTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 176

DB 297 GCGCGGCG 356

QY 177 TCGCCCGAGTCCCGCATAGCGGAGACCTGTACTATTCTCTACCCGCTGTTGTAGG 236

DB 357 GCGTCCCGCGAGGACCGCTCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 416

QY 237 CTTCGCGCGCTGCTGGCTTGGCTGGCGCTTCACCTGGGCGCTCTCTTCTGTGGCTGTGCCA 296

DB 417 GTCCCTGGCGCGCGGACTGTGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 476

GenCore version 5.1.1.6
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CM nucleic - nucleic search, using sw model

Run on: October 24, 2003, 18:12:56 / Search time 1046.7 Seconds
(without alignments)

11633.369 Million cell updates/sec

Title: US-09-830-902-1_COPY_10000_10500

Perfect score: 501

Sequence: 1 ggttcgcgcggtctcgggg.....ggggggggggggggggggcggcggc 501

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 22781192 seqs, 1215238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum Match 100%

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database: EST

1: em_estba:*

2: em_estbm:*

3: em_estbi:*

4: em_estbv:*

5: em_estbo:*

6: em_estbp:*

7: em_estbr:*

8: em_estbs:*

9: gb_esti:*

10: gb_est2:*

11: gb_estc:*

12: gb_est1:*

13: gb_est4:*

14: gb_est5:*

15: em_estfm:*

16: em_estfn:*

17: em_gss_mur:*

18: em_gss_inv:*

19: em_gss_p1:*

20: em_gss_v1:*

21: em_gss_fur:*

22: em_gss_mam:*

23: em_gss_mus:*

24: em_gss_drc:*

25: em_gss_frd:*

26: em_gss_phg:*

27: em_gss_vr1:*

28: gb_gssi:*

29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	344	68.7	916	13 BQ944322	BQ944322 AGENCOURT
2	281.2	56.1	1106	13 BU518568	BU518568 AGENCOURT
3	248.2	49.5	517	28 AQ937943	AQ937943 NB6-679E
4	246	49.1	2897	11 AK044900	AK044900 Mus muscu

RESULT:

BQ944322

LOCUS

DEFINITION

AGENCOURT_8558958 EC1 CGAP Mam2 Mus musculus cDNA clone

IMAGE:6441742 5' mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BQ944322 8558958 EC1 CGAP Mam2 Mus musculus cDNA clone
IMAGE:6441742 5' mRNA sequence.

ACCESSION BQ944322.1 GI:22359600

VERSION BQ944322.1

KEYWORDS EST

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 916)

AUTHORS NIH-VGC <http://mgs.nci.nih.gov/>

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL Unpublished

COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-f@mail.nih.gov
Issue Procurement: Gilbert Smith, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLN);
cDNA sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLN at:
<http://image.lln.gov>
Plate: LLAM-3965 row: g column: 23
High quality sequence stop: 478.

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6	224.8	44.9	600	12	B-987009
7	224.2	44.8	557	13	BM506197
8	220.2	44.0	384	13	BY261238
9	213	43.1	976	10	BQ937372
10	208.8	41.7	553	10	B5618976
11	200.6	40.0	1843	11	AX007193
12	197.6	39.4	537	14	BY707801
13	196.4	39.2	773	14	CB917137
14	188.2	37.6	436	28	AC933698
15	186.6	37.2	548	28	AC933649
16	186.6	37.2	890	10	BF178610
17	183.2	36.6	589	10	B5373511
18	172.8	34.8	569	10	B5625233
19	170.2	34.0	641	12	BM491941
20	164	32.7	604	9	AJ450818
21	163.4	32.6	653	9	AJ446053
22	161.4	28.2	877	9	AL628068
23	161.2	28.2	662	9	AL649334
24	139	27.7	617	9	AL635035
25	138	27.5	623	9	AL655040
26	135.8	27.1	615	9	AL972234
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28	125	25.0	648	9	AL873602
29	124	24.8	644	9	AL871548
30	117.8	23.5	853	13	BQ907712
31	115.8	23.1	683	11	AF549569
32	115	23.0	316	13	BQ571119
33	105.4	21.0	379	14	CJ348493
34	103	20.6	655	10	BB619871
35	99.8	19.9	529	14	C8018332
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43	66.8	13.3	382	13	BX415111
44	66.2	13.2	1057	13	BX496888
45	64.8	12.9	982	13	BX415111

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo; (bases 1 to 517)
 Zabarovsky, E.R., Gizaalil, R., Podowski, R.M., Zabarovsky, V.V., Xie, L., Yuravlenko, O.V., Kozlov, S., Petronko, L., Skoboleva, N., Li, J., Protodov, A., Kishaba, V., Erberg, L., Winberg, G. and Wahlstedt, C.
 Not1 clones in the analysis of the human genome
 Nucleic Acids Res. 28 (1), 1635-1639 (2000)
 20175728
 10710430
 Contract: Podowski RM
 Center for Genomics Research
 Karolinska Institute
 17177 Stockholm, Sweden
 Tel: +46-8-728-6372
 Fax: +46-8-337983
 Email: Raf.Podowski@cr.ki.se
 Class: Not1 site.
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 Best Local Similarity 93.6%; Pred. No. 66-40;
 Matches 278; Conservative 0; Mismatches 17; Indels 2; Gaps 2;
 QY 1 GATTCCCGTCGCTCGCGGCGCGGTTATGCGCGCGCGCGAGCTGTGATG 60
 DB 221 GATTCGCGTCGCTCGCGGCGCGGTTATGCGCGCGCGCGAGCTGTGATG 280
 QY 61 AATTCGCGGCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 170
 DB 281 AATTCGCGGCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 340
 QY 321 CCGAGCGCTCG 190
 DB 341 CCGAGCGCTCG 400
 QY 181 CCGAGCGCTCG 240
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 DB 460 GCGCTCGCGCTCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 516
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 AK044900
 LOCUS
 DEFINITION
 Mus musculus 9.5 days embryo parthenogenote cDNA, RIKEN full-length
 enriched library, clone:B130C10N24 product:spastic paraplegia 4
 homolog (human), full insert sequence.
 AK044900
 AK044900.1 G1:26790554
 HIC; CAP trapper;
 Mus musculus (house mouse)
 Mus musculus
 ORGANISM
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1
 Carninci, P. and Hayashizaki, Y.
 High-efficiency full-length cDNA cloning
 Meth. Enzymol. 303, 19-44 (1999)
 99279253
 PUBLISHED
 10349636
 REFERENCE
 2
 Carninci, P., Shibata, Y., Hayatsu, N., Sugatara, Y., Shibata, K.,

Itch, M., Komno, H., Okazaki, Y., Muramatsu, X. and Hayashizaki, Y.
 Normalization and subtraction of cap-trapper-selected cDNAs to
 prepare full-length cDNA libraries for rapid discovery of new genes
 Genome Res. 10 (10), 1617-1630 (2000)
 20499374
 1042159
 3
 Shibata, K., Itch, M., Azawa, K., Nagacka, S., Sasaki, N., Carninci, P.,
 Komno, H., Akiyama, S., Nishi, K., Kitsuai, T., Tashiro, H., Itch, M.,
 Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
 Yamamoto, S., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
 Fujikake, S., Inoue, K., Togawa, Y., Izawa, M., Chara, E., Watabiki, Y.,
 Yonekura, Y., Shikawa, T., Ozawa, K., Tanaka, T., Matsuda, S., Kawauchi,
 Okazaki, Y., Muramatsu, X., Inoue, Y., Kito, A. and Hayashizaki, Y.
 RIKEN integrated sequence analysis (RISA) system--384-format
 sequencing pipeline with 384 multipillarary sequencer
 Genome Res. 10 (11), 1757-1771 (2000)
 20510913
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 4
 Kawai, J., Shinagawa, A., Shibata, K., Yoshino, M., Itch, M., Ishii, Y.,
 Arakawa, T., Hara, A., Fukunishi, Y., Komno, H., Adachi, S., Fukuda, S.,
 Azawa, K., Izawa, M., Nishi, K., Kiyosawa, H., Kondo, S., Yamanaka, I.,
 Saito, T., Okazaki, Y., Gojohori, T., Bono, H., Kasukawa, T., Saito, R.,
 Kado, K., Matsuda, H., Ashburner, M., Batalov, S., Casavetti, T.,
 Pleschmann, W., Gaasterland, T., Gissi, C., King, L., Kochiwa, H.,
 Kuehl, P., Lewis, S., Matsuo, Y., Nakado, I., Pesole, G.,
 Quackenbush, J., Schrim, L. M., Staub, F., Suzuki, R., Tomita, M.,
 Wagner, L., Washio, T., Sakai, K., Okido, T., Furuno, M., Aono, H.,
 Baldarelli, R., Barsh, G., Blake, J., Boftelli, D., Bolunga, N.,
 Carninci, P., de Bona, M. F., Brownstein, M., Bult, C.,
 Fletcher, C., Fujita, M., Gariboldi, V., Gustincich, S., Hill, D.,
 Hofmann, M., Hume, D. A., Kamiya, M., Lee, N. H., Lyons, F.,
 Marchionni, L., Mashima, J., Mazzarelli, J., Mombarts, P., Nordone, P.,
 Ring, B., Ringwald, M., Rodriguez, I., Sakamoto, N., Sasaki, H.,
 Sato, K., Schonbach, C., Seya, T., Shibata, Y., Storch, K. P., Suzuki, H.,
 Toyooka, K., Wang, K. H., Weitz, C., Whittaker, C., Wilming, L.,
 Wynshaw-Boris, A., Yoshida, K., Hasegawa, Y., Kawaji, H., Kohtsuki, S.,
 and Hayashizaki, Y.
 Functional annotation of a full-length mouse cDNA collection
 Nature 409 (6821), 695-690 (2001)
 21085660
 11217851
 5
 The PANTON Consortium and the RIKEN Genome Exploration Research
 Group Phase I & II Team.
 Analysis of the mouse transcriptome based on functional annotation:
 of 40,778 full-length cDNAs
 Nature 420, 563-573 (2002)
 6 (bases 1 to 2897)
 Adachi, J., Arakawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
 Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
 Hayashida, K., Hayatsu, N., Hiramoto, K., Hirata, T., Hirozane, T.,
 Horii, P., Imotani, K., Ishii, Y., Itch, M., Kagawa, I., Kasukawa, T.,
 Kato, H., Kawai, S., Kojima, Y., Kondo, S., Komno, H., Kouda, M.,
 Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M.,
 Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N.,
 Okazaki, Y., Saito, K., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N.,
 Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
 Sugabe, Y., Tamai, M., Tagawa, A., Takanashi, F., Takaku-Akanishi, S.,
 Takada, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A.,
 Muramatsu, M. and Hayashizaki, Y.
 Direct Submission
 Submitted (16-JUL-2001) Yoshitake Hayashizaki, The Institute of
 Physical and Chemical Research (RIKEN), Laboratory for Genome
 Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
 RIKEN Yokohama Institute, 1-7-22 Sueno-cho, Tsurumi-ku, Yokohama,
 Kanagawa 230-0045, Japan (E-mail: genome-res@gsc.riken.go.jp,
 URL: http://genome.gsc.riken.go.jp/, Tel: 81-45-503-9222,
 Fax: 81-45-503-9216)
 CDNA library was prepared and sequenced in Mouse Genome
 Encyclopedia Project of Genome Exploration Research Group in Riken
 Genomic Sciences Center and Genome Science Laboratory in RIKEN.
 COMMENT


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RESULT 6
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VERSION B1987009.1 GI:7957970
KEYWORDS EST.
SOURCE Mus musculus (house mouse).
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 600)
AUTHORS Mu, X., Zhao, S., Pershad, R., Hsieh, T. F., Scarpa, A., Wang, S. W.,
White, R. A., Beremand, P. F., Thomas, T. J., Gan, H., and Klein, W. H.
TITLE Gene expression in the developing mouse retina by EST sequencing
and microarray analysis
JOURNAL Nucleic Acids Res. 29 (24): 4963-4993 (2001)
MEDLINE 21671925
PUBMED 11812828
COMMENT Contact: Klein W.
Department of Biochemistry and Molecular Biology
University of Texas M.D. Anderson Cancer Center
Box 117, 1515 Holcombe Blvd., Houston, TX 77030, USA
Tel: 713 792 3646
Fax: 713 792 0329.
Location/Qualifiers
organism="Mus musculus"
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tissue_type="neural retina"
dev_stage="embryonic day 14.5 post-fertilization"
clone_lib="Mouse E14.5 retina la-bda zap II library"
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 114305.y1 Melton: Normalized Mixed Mouse Pancreas: NI-MMS1 MMS
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ACCESSION: BMS06397
 VERSION: BMS06397.1 GI:18670439
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 SOURCE: Mus musculus (house mouse).
 ORGANISM: Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 REFERENCE 1 (bases 1 to 557)
 AUTHORS: Melton, D., Brown, J., Kent, J., Permut, A., Lee, C., Kaestner, K.,
 Lerishka, I., Secor, M., Brestelli, J., Gradwohl, G., Clifton, S.,
 Hallier, L., Warr, M., Page, D., Wylie, T., Martin, J., Bistain, A.,
 Schmitt, A., Theising, B., Rutter, E., Ronko, J., Bennett, L., Cardenas,
 X., Gibbons, M., McCann, R., Cole, R., Tsagaris, R., Williams, T.,
 Jackson, Y., and Bowers, Y.
 TITLE: Endocrine Pancreas Consortium
 JOURNAL: Unpublished
 COMMENT: Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue
 Endocrine Pancreas Consortium
 Harvard University, Howard Hughes Medical Institute
 Dept. of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
 MA 02138
 Tel: 617-495-1812
 Fax: 617-495-8557
 E-mail: dmelton@biochem.harvard.edu
 Library was constructed by Dr. Douglas Melton. DNA sequencing by:
 Washington University Genome Sequencing Center. For information on
 obtaining a clone please contact: Juliana Brown
 jbrown@wustl.edu
 NCBI:207249 This sequence now available from the IMAGE consortium,
 for clone orders contact: info@image.llnl.gov
 Seq primer: -45RP from Gibco
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 Location/Qualifiers
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 islets of Langerhans were separately constructed using
 Superscript Plasmid Library kit (Life Technologies). cDNA
 was made by oligo-dT priming and size-selected by column
 fractionation. Libraries were amplified once on solid

support and plasmid DNA from each library was prepared and mixed in equal amounts. The mixed library DNA was normalized by method #4 from Benaldo, Lennon, and Soares 1996 Genome Research 6:791-806; 5.5 microgram single-stranded mixed library plasmid DNA was mixed with 5 micrograms PCR product representing mixed library inserts and hybridized to an EcoT of 6. Single-stranded (unhybridized) plasmids were isolated by hydroxyapatite chromatography and used to make this library."

BASE COUNT 132 a 164 c 152 g 109 t
ORIGIN

Query Match 44.8%; Score 224.2; DB 12; Length 557;
Best Local Similarity 89.1%; Pred. No. 3.8e-15;
Matches 244; Conservative 0; Mismatches 33; Indels 0; Gaps 0;

QY 196 AAGCGGACCTGTACTATCTCTACCCCTCTTTGTAGCTTCGGCTGCTGCTG 255
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RESULT 8
BY261238
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BY261238 384 bp mRNA linear EST 10-DEC-2002
BY261238 RIKEN full-length enriched, visual cortex Mus musculus
cDNA clone K33035P2 5', mRNA sequence.

BY261238
BY261238.1 GI:26442750
EST
Mus musculus (house mouse)

Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sclerognathi; Muridae; Murinae; Mus.

1 (bases 1 to 384)
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Enno, H., Kondo, S.,
Nikaido, T., Osato, N., Saito, R., Suzuki, H., Yamahara, Y., Kiyosawa, H.,
Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schorbach, C.,
Gojoh, T., Baidar, R., Hill, D. P., Bitt, C., Hume, D. A.,
Quackenbush, J., Schirf, J. V., Kanapin, A., Matsuda, H., Batalov, S.,
Beisel, K. W., Blake, J. A., Brad, D., Brusis, V., Cho, H. A., Corbett, L. E., Cousins, S., Datta, E., Dragani, T. A., Fletcher, C. E., Forrest, A., Frazer, K. S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, C., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I. J., Garvis, E. D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R. Y., King, B. L., Konagaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons, P. A., Maglott, D. R., Maltais, D., Marchionni, J., McKenzie, L., Mik, H., Nagashima, T., Numata, K., Okido, T., Pavan, W. J., Petosa, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J. U., Q. D., Ramachandran, S., Ravasi, T., Reed, J. C., Reid, J. C., Reid, J. C., Ring, B. Z., Ringwald, M., Sander, A. A., Schneider, C., Seiple, A., Setzu, M., Shimada, K., Sultana, R., Takanaka, Y., Taylor, N. S., Teasdale, R. C., Tomita, M., Verardo, R., Wagner, L., Wamboldt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagisawa, Y., Yang, L., Yang, L., Yuan, Z., Zavolan, M., Zhu, X., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, G., Arakawa, K., Arakawa, T., Fukuda, S., Hara, A., Hashizume, N., Imotani, K., Ishii,

Y., Itoh, M., Kacawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E. S., Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
22354683
PUBMED
12466851

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Fax: 81-45-503-9216
Email: genome-resgsc.riken.go.jp
URL: http://genome.resc.riken.go.jp/

Arakawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numata, K., Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tatem, M., Waki, K., Watanabe, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission
Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome 12, 673-677 (2001).
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN integrated sequence analysis (RISA) system-384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1751-1771 (2000)
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in Riken Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.
Tissues were provided by Michela Fagioli and Takao K. Hersch, Laboratory for Neuronal Circuit Development Brain Science Institute RIKEN 2-1, Hirotsawa, Wako-shi, Saitama 351-0198 Japan, whose assistance we gratefully acknowledge. Please visit our web site at: http://genome.resc.riken.go.jp/ for further details.

Location/Qualifiers
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Matches 243; Conservative 0; Mismatches 35; Indels 0; Gaps 0;

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ACCESSION BY707801.1 GI:2711898;
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KEYWORDS EST.
SOURCE Mus musculus (house mouse);
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REFERENCE
AUTHORS

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1. (bases 1 to 597)
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nakado, I., Otsu, N., Saito, R., Suzuki, R., Yamakawa, I., Kiyosawa, H.,
Yagi, K., Toraru, Y., Hasegawa, Y., Noga, A., Schorbach, C.,
Gocho, T., Baldarelli, R., Hill, D.P., Bult, C., Hume, D.A.,
Quackenbush, J., Schram, L.M., Kanapir, A., Matsuda, H., Batalov, S.,
Reisel, K.W., Blake, J.A., Bradt, D., Brusic, V., Cho, T.A., Corbaci,
L.E., Cousins, S., Dalla, E., Dragani, T.A., Fletcher, C.F., Forrest,
A., Frazer, K.S., Gaasterland, T., Gaibolli, M., Gissi, C., Godzik, A.,
Gough, J., Grigoriadis, S., Gustincich, S., Hirokawa, N., Jackson, I.,
Karsis, E.D., Kanai, A., Kawaji, R., Kawasawa, Y., Kedzierski, R.M.,
King, B.L., Konagaya, A., Kurochkin, I.V., Lee, Y., Lenhard, B., Lynch,
P.A., Maglott, D.R., Maltsev, K., Marchionni, L., McKenzie, L., Miki,
H., Nagasima, T., Numata, K., Okado, T., Pavan, W.J., Petosa, C.,
Pescio, G., Petrovsky, N., Pilla, R., Pontius, J.J., Qian, Q.,
Ramachandran, S., Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring,
B.Z., Ringwald, M., Sandelin, A., Schneider, C., Sepp, C.A., Setou,
M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, X.S., Teasdale,
R.D., Tomata, Y., Verardo, R., Wagner, L., Wastle, C., Wang, Y.,
Watanabe, Y., Wells, C., Wilming, L.G., Wynshaw-Boris, A., Yanagisawa,
M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A.,
Carninci, P., Hayashino, N., Hirozane-Kishikawa, T., Kono, H., Nakamura,
Y., Sakazume, N., Sato, K., Shiraki, T., Wak, K., Kawai, J., Azakura, K.,
Aizawa, T., Fukuda, S., Harai, A., Hashizume, W., Iotani, K., Ishii,
Y., Itoh, K., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, C., Shibata,
K., Shiragawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Zander,
E.S., Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
22354683
2466851
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
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The Institute of Physical and Chemical Research (RIKEN)
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Tel.: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome.res@gs.c.riken.go.jp/
URL: http://genome.gsc.riken.go.jp/
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda,
S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F., Imotani, K.,
Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y., Kondo, S., Kono,
H., Koya, S., Miyazaki, A., Murata, M., Nakamura, Y., Nomura, K.,
Numazaki, R., Ohsato, N., Saito, R., Sakazume, N., Sato, H.,
Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y.,
Waki, K., Watanabe, A., Murakami, M. and Hayashizaki, Y. Direct
Submission
Computational Analysis of Full-length Mouse cDNAs Compared with
Human Genome Sequences. Yarn. Genome 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
Genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN Integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer. Genome Res.
10 (11), 1757-1771 (2000)
Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
redundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Please visit our web site (http://genome.gsc.riken.go.jp) for
further details.
Location/Qualifiers
1. 597
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GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OX nucleic - nucleic search, using sw model

Run on: October 24, 2003, 15:43:05 : Search time 13el.12 seconds
(without alignment)
14808.112 Million cell updates/sec

Title: US-09-830-902-1_COPY_1_500

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Scoring table: IDENTITY_NUC

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Searched: 2655711 seqs, 20454815386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0

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Post-processing: Minimum Match 0%

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Listing first 45 summaries

Database :

GenEmbl:

1: gb_bai*

2: gb_htg*

3: gb_in*

4: gb_em*

5: gb_ov*

6: gb_pat*

7: gb_ph*

8: gb_pl*

9: gb_dr*

10: gb_ro*

11: gb_sts*

12: gb_sy*

13: gb_un*

14: gb_vl*

15: em_ba*

16: em_fun*

17: em_hum*

18: em_ir*

19: em_mu*

20: em_or*

21: em_ov*

22: em_pat*

23: em_ph*

24: em_pi*

25: em_ro*

26: em_sts*

27: em_un*

28: em_vl*

29: em_vl*

30: em_htg_hum*

31: em_htg_inr*

32: em_htg_other*

33: em_htg_mus*

34: em_htg_pln*

35: em_htg_rod*

36: em_htg_mar*

37: em_htg_vrt*

38: em_sy*

39: em_htgo_hum*

40: em_htgo_mus*

41: em_htgo_other*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	% Match	Query Length	DB ID	Description
1	500	100.0	101584	9	CNS01DSS
2	500	100.0	110000	6	AL121655 BAC seque
3	500	100.0	110000	6	AX093471 Sequence
4	500	100.0	110000	6	AX246003 Homo sapi
5	231.8	40.8	139568	9	AC012364 Homo sapi
6	180.8	36.2	117751	9	HS16915 Homo DNA s
7	189.6	38.0	45195	9	AC023913 Homo sapi
8	189.6	37.9	156589	9	AC090421 Homo sapi
9	187.6	37.5	172837	9	HS537K23 Human DNA
10	181.4	36.3	169334	9	AL453304 Human DNA
11	181.4	36.3	18434	2	AC135817 Papio anu
12	179.4	35.9	59329	2	AC135815 Papio anu
13	179.4	35.9	57621	9	AL929255 Homo sapi
14	179.4	35.9	121362	9	AL607089 Human DNA
15	179.4	35.9	145736	9	AL359203 Human DNA
16	179.4	35.9	152428	9	AL451139 Human DNA
17	179.2	35.8	134337	9	AC114492 Homo sapi
18	179	35.8	15923	9	HSB436C9 Human DNA
19	177.4	35.5	188546	2	AC01504 Pan trogl
20	177.2	35.4	171309	9	AC023396 Homo sapi
21	176.8	35.4	90214	9	AC012146 Homo sapi
22	176.8	35.4	190649	9	AL356488 Human DNA
23	176.6	35.3	186902	9	CNS01DX1 Human chr
24	176.4	35.3	90243	9	AC010863 Homo sapi
25	176.4	35.3	151980	9	AL669876 Human DNA
26	176	35.2	106034	9	AC084209 Homo sapi
27	176	35.2	154788	2	AC011984 Homo sapi
28	175.8	35.2	186252	2	AC144518 Pan trogl
29	175.8	35.2	110002	9	AE014305_0 Homo sapi
30	175.8	35.2	18406	9	AL445188 Human DNA
31	175.6	35.1	204439	9	AF003637 Homo sapi
32	175.6	35.1	39729	6	AX647093 Sequence
33	175.6	35.1	64475	6	AF283320S2 Homo sapi
34	175.6	35.1	66933	6	AX277532 Sequence
35	175.6	35.1	66933	6	AX418096 Sequence
36	175.6	35.1	137206	2	AC024123 Homo sapi
37	175.6	35.1	159720	2	AF023366 Homo sapi
38	175.4	35.1	75167	9	AC092029 Homo sapi
39	175	35.0	185087	9	AC093884 Homo sapi
40	175	35.0	128152	9	AL136313 Human DNA
41	174.8	35.0	59257	9	AC106019 Homo sapi
42	174.8	35.0	110144	9	AC080053 Homo sapi
43	174.8	35.0	130855	9	AC084069 Homo sapi
44	174.8	35.0	188340	9	AC135178 Homo sapi
45	174.8	35.0	189623	2	AC018640 Homo sapi

ALIGNMENTS

RESULT 1
CNS01DSS
LOCUS
DEFINITION BAC sequence from the SP04 candidate region at 2p21-2p22 BAC 336P14
of library CITB_978_SKB from chromosome 2 of Homo sapiens (Human).
ACCESSION AL121655
VERSION AL121655.1 GI:6002386
KEYWORDS SP04 genomic DNA interval.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 101584)
AUTHORS Hazan,J., Fonknechten,N., Mavel,D., Paretnette,C., Sanson,D.,
Artiguenave,F., Davoine,C.S., Cruaud,C., Durr,A., Wincker,P.,


```

QY 301 TCCTCTGCTCAACTCTTCCAACTAGCTGGGACTACAGAAATGACTCCGCTGCTGAC 340
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QY 421 ACTTTGGGAGACCAAGGCGGCTACTTACCTAGGTCAGAGCTTCAAGCAGCAGCGCC 460
Db 421 ACTTTGGGAGACCAAGGCGGCTACTTACCTAGGTCAGAGCTTCAAGCAGCAGCGCC 460
QY 481 AACATGGTAAACCATGTGCG 500
Db 481 AACATGGTAAACCATGTGCG 500

RESULT 3
LOCUS HSA246003 110000 bp DNA linear FBI 08-MAR-2000
DEFINITION Homo sapiens Spast gene for spastin protein.
ACCESSION AC245003
VERSION AC246003.1 G:6273492
KEYWORDS Spast gene; spastin protein; SPG4-linked hereditary spastic paraplegia.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Hazan, J., Fonknechten, N., Mavel, D., Paternotte, C., Samson, D., Artiguenave, F., Davoine, C.S., Cruaud, C., Durr, A., Winkler, P., Brodier, P., Cattolico, L., Barbe, V., Burgunder, J.M., Prud'homme, J.F., Brice, A., Fontaine, B., Heilig, R., and Weissbach, J.
TITLE Spastin, a new AAA protein, is altered in the most frequent form of autosomal dominant spastic paraplegia
JOURNAL Nat. Genet. 23 (3), 296-303 (1999).
MEDLINE 20055425
PUBMED 10610178
REFERENCE 2
AUTHORS Fonknechten, N., Mavel, D., Byrne, P., Davoine, C., Cruaud, C., Boetsch, D., Samson, D., Costinbo, P., Hutchinson, M., McMonagle, P., Burgunder, J., Tarragione, A., Heitzel, G., Rekt, I., Douflet, T., Parfey, N., Brice, A., Fontaine, B., Prud'homme, J., Weissbach, J., Durr, A., and Hazan, J.
TITLE Spectrum of SPG4 mutations in autosomal dominant spastic paraplegia
JOURNAL Hum. Mol. Genet. 9 (4), 637-644 (2000)
MEDLINE 20164102
PUBMED 10693187
REFERENCE 3 (bases 1 to 110000)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (17-JUN-1993) Genoscope, Genoscope - Centre National de Sequencage, BP 191, EVRY 91006, FRANCE
COMMENT E-mail : seqraf@genoscope.cns.fr - web : www.genoscope.cns.fr. The sequence is the result of the assembly of 2 BAC clones: P-336p.4 and 563N4, respectively from RP11-II and CtrB_SPS_SKB library.
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73249..74613
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gene

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an accurate chemistry, or covered by high quality data (i.e., phred quality > 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/jsc>

SOURCE INFORMATION:

The RPCL11 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Woon, P.Y., Zhao, S., Freeman, E., Tateo, M., Catanese, J.J. and de Jong, P.J. (1995). An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 5:11-9. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RPCL11-41p13; the clone sequenced to the right is RPCL11-78p13. Actual start of this clone is at base position 1 of RPCL11-44p15; actual end is at base position 155943 of RPCL11-44p15.

Data from AC011232 and AC010981 was used to finish this clone. AC012364. The sequence fidelity between bases 112626 to 112643 can not be guaranteed due to an unresolved homopolymeric run. The sequence between 112648 to 112663 is single stranded.

FEATURES

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Matches 500; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 115737 TAACTGACTGCTGCTGTTTCCTTGTCATATATAGCTAATCAATGAGCAGGTCAAGTAA 115796
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CY 121 AACAAATCCCTCCATACACTAGAAATCTCAAACTGGTGAATGGGAGTTTGGTTTGT 180
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DB 115857 AACAAATCCCTCCATACACTAGAAATCTCAAACTGGTGAATGGGAGTTTGGTTTGT 115916
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CY 181 TTTTGTTGTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTT 240
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DB 116157 ACTTTGGAGACCAAGCCAGCCGCAATTTACTTGAAGTTAGAGTTCAGACCAAGCCAGGCC 116216
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CY 481 AACATGGTAAACCATCTCG 500
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DB 116217 AACATGGTAAACCATCTCG 116236
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RESULT 5
HS16915/c 133568 bp DNA linear PRI 13-NOV-2000
LOCUS Human DNA sequence from clone RPI-16915 on chromosome Xp22.
DEFINITION Contains ESTs, STSs, GSSs and CpG islands. Contains the 3' part of
the DDX3 gene for DEAD/H (Asp-Glu-Ala-Asp/His) box polypeptide 3
and a novel gene for a leucine rich protein, complete sequence.
ACCESSION Z93015
VERSION Z93015.9 GI:9929207
KEYWORDS HTG; CpG is.; and; DDX3; DEAD box; leucine rich protein.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 133568)
AUTHORS Pavitt,R.
DIRECT SUBMISSION
TITLE Submitted (24-OCT-2000) Sanger Centre, Hinxton, Cambridgeshire,
C81C 1SA, UK. E-mail: enquiries@humquerry@sanger.ac.uk
requests: clonerquest@sanger.ac.uk
COMMENT On Aug 26, 2000 this sequence version replaced gi:9798426.
```

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'uncure' feature key.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/ChrX>

RPI-16915 is from the library RPI-1 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffa.edu/> VECTOR: pCYPAC2

This sequence is the entire insert of clone RPI-16915.

Location/Qualifiers

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/evidence="not_experimental"

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/note="match: STS: Em:G34988"

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/gene="DDX3"

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/gene="DDX3"

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/note="L1MC5 repeat: matches 1741..7890 of consensus"

3392..3431

/note="L1MC5 repeat: matches 5776..5817 of consensus"

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/note="L1MC5 repeat: matches 5703..5776 of consensus"

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/note="L1MC5 repeat: matches 7933..7916 of consensus"

3944..4108

/note="MER5B repeat: matches 1..161 of consensus"

5253..5783

/note="L1MC5 repeat: matches 5669..6163 of consensus"

6086..6477

/note="L1MC5 repeat: matches 5182..5669 of consensus"

6778..6881

/note="L1MC5 repeat: matches 5286..5182 of consensus"

7186..7530

/note="L1 repeat: matches 3143..3507 of consensus"

7854..8044


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Tr:Q61332 Tr:Q95123 Sw:P10075 Tr:Q9401 Sw:P17142
Tr:Q9CUN7 Tr:Q9Y545 Sw:P15523 Tr:Q60304 Sw:P10754
Tr:Q62517 Tr:Q00146 Sw:P18713 Tr:Q94JW8
Tr:Q62576 Tr:Q9W35 Tr:Q9Y2P1 Sw:Q62513 Tr:Q90658
Sw:P52738 Tr:Q9W35 Tr:Q94178 Sw:P24358 Wp:C5587 Tr:Q9VD05
Sw:Q95125 Tr:Q9UP97 Sw:P18724 Sw:P15613 Tr:Q57415
Tr:Q9WVJ8 Sw:P18729 Tr:Q62886 Tr:Q913M9 Tr:Q9Y467
Wp:Y55FJAM Tr:Q9R161 Sw:P18747 Tr:Q9JW8 Sw:P18749
Tr:Q74256 Sw:Q14628 Sw:Q02386 Tr:Q95878 Sw:P18736
Sw:P18738"
/pseudocoding
/codon_start=1
/revidence=not_experimental
41524..41878
/gene="dJ537K23.2"
/note="match: GSS: Em:AQ341439"
/pseudocoding
repeat_region 42823..43053 /note="MER41-internal repeat: matches 2967..3199 of
consensus"
repeat_region 43352..43429 /note="MER41-internal repeat: matches 3199..3275 of
consensus"
repeat_region 43683..44198 /note="MER41-internal repeat: matches 3275..3941 of
consensus"
44241..46953
/note="ITR1 repeat: matches 1..785 of consensus"
47857..48721
/note="LIP3 repeat: matches 5293..6100 of consensus"
48833..48920
/note="26 copies 2 mer aa 73% conserved"
52743..52886
/note="LIM4 repeat: matches 6442..6845 of consensus"
57849..57901
/note="AluJ/monomer repeat: matches 233..302 of consensus"
58354..58741
/note="H5E4 repeat: matches 1..371 of consensus"
63307..63348
/note="21 copies 2 mer tt 78% conserved"
63539..63588
/note="S:nole clone region"
65820..65923
/note="AluJ/F(R)AM repeat: matches 149..297 of consensus"
66814..67015
/note="VER63A repeat: matches 1..210 of consensus"
71827..71896
/note="2 copies 35 mer 90% conserved"
75291..75714
/note="match: GSS: Em:AQ72817"
75766..76167
```

```

of Plaster de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACE1.6
This sequence is the entire insert of clone RP11-505Q17. The true
left end of clone RP11-18k13 is at 14759; in this sequence, the
true right end of clone RP11-333G16 is at 113781 in this sequence.
FEATURES             Location/Qualifiers
     source            1..172937
                     /organism="Homo sapiens"
                     /mol_type="genomic DNA"
                     /db_xref="taxon:9606"
                     /chromosome="1"
                     /clone_lib="RP11-505Q17"
                     /clone_name="RP11-333G16"
misc_feature          18441..18559
                     /note="Sequence from overlapping clone RP11-333G16
                     (AL590620). Assembly confirmed by restriction digest."
misc_feature          22613..22657
                     /note="Sequence from overlapping clone RP11-333G16
                     (AL590620). Assembly confirmed by restriction digest."
misc_feature          24337..26486
                     /note="Sequence from overlapping clone RP11-333G16
                     (AL590620). Assembly confirmed by restriction digest."
misc_feature          93691..100086
                     /note="Single clone region. Assembly confirmed by
                     restriction digest data."
misc_feature          99946..100202
                     /note="Sequence from uni-directional cDNA big dye
                     terminator reads only."
misc_feature          100235..100319
                     /note="Sequence from overlapping clone RP11-333G16
                     (AL590620). Assembly confirmed by restriction digest."
misc_feature          127409..127498
                     /note="Single clone region. Sequence from reads from a
                     short insert library derived from a single p3C clone.
                     Restriction digest data confirm the assembly."
BASE COUNT           49503 a 33911 c 34461 g 54962 t
ORIGIN
Query Match      37.5%; Score 187.6; DB ?; Length 172937;
Rest Local Similarity 75.2%; Pval No. 2.5e-17;
Matches 248; Conservative 0; Mismatches 79; Indels 3; Gaps 17
QY   170 TGGTTTGTGTTTTGTTTCTATTATTTATTTATTTATTTATTTAGATGGAGTTTGCT 228
Db    43942 TGAGTTCCCAATTTGGTGTGTTTGGTGTGTTTGGTGTGTTTGAAGAAGAGTTCAC 44301
QY   220 CTGTTGCCAGGCGTGGAAATGCATGCATGCATGCATGCATGCATGCATGCATGCATGC 269
Db    44002 CTGTTGCCAGGCGTAGGCGCATGCTGCAGTCTCAGCTCAGCTCAGCTCAGCTC 44561
QY   290 GGTTCAAGCAATCCTGCTGCTCAATCTTCCAAAGTAGCTGGCAGTACAGGAATGAGCTGC 349
Db    44062 GGTTCGAAGCAATTTCTCATGCCCTCAGACTCCACGTAGTTGGGATTTACAGGCATGAGACAC 44121
QY   350 CGAGCTGGCGCTGGTTGTTGTTTGTATTAATTTTGAAGCGAGGTGCAGTGGCGGCTATCT 409
Db    44122 CAGCTTGGCCAG---TTATTTTATAAAGTTTTTGGCCAGGCGAGGTGGGCTCAGGCT 44178
QY   410 GTGATCCCAGCACATTTGGGAGACAAGGAGGCGCGGCGGATTACTTGAAGTCAGGAGTCAAGA 469
Db    44179 GTAATCCCAGCACATTTGGGAGGACAAGGCGAGGTGATCACTTGAGGTCAAGAGTTTGACA 44238
QY   470 CCAGCCAGGCGCACATGGTAAAACCATGTC 499
Db    44239 CCAGTTTGGCCAACATGTTCAAACCCCATC 44268

```

```

RESULT 10
AC135837
LOCUS      AC135837      169934 bp      DNA      linear      HTG 16-JAN-2003
DEFINITION Papio anubis clone RP41-316H21, WORKING DRAFT SEQUENCE, 6 ordered
            pieces.

```

AC135837
VERSION
G:127502021
HTG: HTGS PHASE2, HTGS DRAFT.
SOURCE
Papio anubis (olive baboon)
ORGANISM
Papio anubis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea; Cercopithecinae; Papio.
1 (bases 1 to 169934)
AUTHORS
Akhtar, N., Antonellis, A., Ayale, K., Beckstrom-Sternberg, S.M., Ben-Amin, B., Brakesley, R.W., Boufard, G.G., Brinkley, C., Brock, C., Carraga, K., Coleman, B., Engle, J., Granite, S., Guan, X., Gupta, J., Haghighi, P., Han, J., Hansen, N., Ho, S.-L., Idci, J.R., Karlins, E., Latic, P., Lee-Lin, S.-C., Legaspi, R., Maduro, O.L., Maduro, V.B., Pargulies, E.H., Mastello, C., Maskeri, B., McDowell, J., Parguiran, C., Pearson, R., Portnoy, M.E., Prasad, A., Reddy-Dugue, N., Schandler, K., Schueler, M.G., Sison, C., Statirpops, S., Thomas, J.W., Thomas, P.J., Touchman, J.W., Vogt, J., Wetberly, K.D., Wiggins, L., Young, A. and Green, E.D.
NISC Comparative Sequencing Initiative
Unpublished
2 (bases 1 to 169934)
AUTHORS
Green, E.D.
TITLE
Direct Submission
JOURNAL
Submitted (23-OCT-2002) NIH Intramural Sequencing Center, 8717 Groveomt Circle, Gaithersburg, MD 20877, USA
3 (bases 1 to 169934)
REFERENCE
Green, E.D.
AUTHORS
Direct Submission
TITLE
Submitted (18-JAN-2003) NIH Intramural Sequencing Center, 8717 Groveomt Circle, Gaithersburg, MD 20877, USA
JOURNAL
On Jan 18, 2003 this sequence version replaced gi:27261523.
COMMENT
----- Genome Center
Center: NIH Intramural Sequencing Center
Center code: NISC
Web site: <http://www.nisc.nih.gov>
Contact: nisc_rcc@nhgri.nih.gov
----- Project Information
Center project name: c9q
Center clone name: 316H21

The sequence data in this record represents an 'enhanced' version of a Phase 2 submission. Specifically, the indicated order and orientation of each sequence contig has been established using one or more of the following: read-pair data from individual sub-clones, overlaps with neighboring clones, alignment with available reference sequence (e.g., human), and/or confirmation by PCR testing. In addition, the sequence assembly is based on at least 8X average coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

..... Summary Statistics

Summary Statistics

Sequencing vector: plasmid; n/a; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990.19
Consensus quality: 16911 bases at least Q40
Consensus quality: 169324 bases at least Q33
Consensus quality: 169383 bases at least Q20
Insert size: 16300; agarose-ff
Insert size: 169434; sum-of-connigs
Quality coverage: 1.81x in Q20 bases; agarose-ff
Quality coverage: 1.16x in Q20 bases; sum-of-connigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* been provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and

- * is believed to be correct as given, however the sizes
- * of the gaps between them are based on estimates that have
- * provided by the submitter.

* This sequence will be replaced
by the sequence shown below.

- by the finished sequence as soon as it is available and

- the accession number will be preserved.

1 5018: contig of 5018 bp in length

* 5019 5118: gap of unknown length
5019 5119: contig of 5500 bp in length

- 60119: consig of 5500, 69 in target.
- 60125: gap of unknown length
- 60126: gap of unknown length

60129 gap of unknown length
60226 contig of 7125 bp length

* 67351 67450: gap of unknown length

* 67451 109121: contig of 41671 bp in length

```

:09:22 :09221: gap of unknown length
:09:22 :09221: gap of unknown length

```

- 109222 247612: contig of 38391 bp in length

* 147623 147712: gap of unknown length.
* 147733 150381: composed of 2669 bp in 5' end +

147713 150381: Schlegel of 2669 pp 1: English
150382 150491: gap of unknown length

* 150482 150481: gap of unknown length
150480 17:241: contig of 20763 bp in length

- 171242 172341: gap of unknown length

* 171342 184341: contig of 1300 bp in length.

 Location/Qualifiers |

184341

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/organism="E. coli"
/mol_type="genomic DNA"
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/mot type=genomic DNA
/db xref="taxon:955"

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/clone="RP41-232D"
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/clone_lib="RP41"
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Signature _____
 1. 45592 _____

/note="Clone overlaps with: GenBank Accession Number NC135547, Clone 3341-2.3P8 (center), protect, name, cco

Signature 1 5012
AC135547 clone RFL-2.3D9 (GenBank project name seq)

[illegible]

```

7...000= 0000 0000 _arguments
clone end:77

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This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digests. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Emi, EMBL; Swi, SWISSPROT; Trl, TREMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep. This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr1>. RP11-547P4 is from the library RP11-11.2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>.

FEATURES
source
Location/Qualifiers
1..121362
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="1"
/clone="RP11-547P4"
/clone_lib="RP11-11.2"
BASE COUNT 36463 a 23081 c 24157 g 37661 t
ORIGIN

Query Match 35.9%, Score 179.4; DB 9; Length 121362;
Best Local Similarity 76.4%; Pred. No. 8.6e-35;
Matches 233; Conservative 0; Mismatches 71; Indels 1; Gaps 1;
QY 195 TTTATTTTATTTTATTTTCTAGATGGAGTCTCTCTGTGTCGCCAGCTGGATGCAATG 254
Db 90665 TTTTCTTTTCTTTTCTTTTCTAGACAGATCTACTCTGGGCGGAGCTGGGATG 90626
QY 255 GCATGATCAGTCTACTGACCTGACCTTTCGGGTTCAAGCAATCTCTGCTCAAA 314
Db 90605 GCATGATCGGGCTACTGCACTCCGGCTCCGGGGCTCAATGATTCCTGCTCAAG 90746
QY 315 TCTTCCAGTAGCTGGATACAGGATGAGTGGCGGACCTGGCTGCTTCTTCTT 374
Db 90745 CTCCCATGTAGTTAGATTACAGGCGCCGACACACACATTTGCGTCCCATATTTTA 90684
QY 375 TTAATTTTGGAGCCAGTGCAGTGGCCCATATCTGTGATTCGACGACCTTGTGAGCA 434
Db 90685 AGAGATAGT-TGGCCTGGTGTAGTACTCATGCTGTATTCACAGCACCTTGGAGGCG 90627
QY 435 AGGACGCGGATTTACTTGGGTCGAGGTCGACAGCAGCGGACGCAATGTTAAACC 494
Db 90626 AGCAGGCGGATTTACCTGAGGTAGAGTTGCGAGCAGCGCTGGACACCGGTGAACC 90567
QY 495 ATGTC 499
Db 90566 CAGTC 90562

RESULT 15
AL451139 145736 bp DNA linear PR: 22-MAY-2002
LOCUS
DEFINITION Human DNA sequence from clone RP11-569G9 on chromosome 1, complete
sequence.
ACCESSION AL451139
VERSION AL451139.40 GI:121211745
KEYWORDS HTG
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 145736)

AUTHORS TITLE JOURNAL

Van Heilmond, Z.
Direct Submission
Submitted (21-MAY-2002); Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail: enquiries@
sanger.ac.uk
 enquiries@sanger.ac.uk
 Clone requests: clonerequests@sanger.ac.uk
 On May 25, 2002 this sequence version replaced gi:20973434.

COMMENT

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Emi, EMBL; Swi, SWISSPROT; Trl, TREMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep. This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr1>. RP11-569G9 is from the library RP11-11.2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>.

FEATURES
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Location/Qualifiers
1..145736
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="1"
/clone="RP11-569G9"
/clone_lib="RP11-11.2"
BASE COUNT 36953 a 37564 c 36509 g 34710 t
ORIGIN

Query Match 35.9%, Score 179.4; DB 9; Length 145736;
Best Local Similarity 75.1%; Pred. No. 8.6e-35;
Matches 238; Conservative 0; Mismatches 76; Indels 3; Gaps 1;
QY 186 TTGTTTATTTTATTTTATTTTATTTTCTAGATGGAGTCTTGTCTGTGTCGCCAGGTGG 245
Db 122735 TTGAAGCACTTTTCTTTCTTTTGTGATGGAGTCTTGTCTGTGTCGCCAGGTGG 122854
QY 246 AATGCAATGATGATCTCAGCTCAGTCTGCAACCTTCCGCTTTCGGGTTCAAGCAATGCTC 305
Db 122855 AGTSCAATGGCGTATTTTGTACTCAGCGCAACCTTCTGCTCTGGGTTTCGAGGATTTCT 122814
QY 306 CTGCTCAATCTTCCAAAGTACTGGGACTACAGGAATGAGTGGCGGACCTGGCTGGT 365
Db 122915 CTGCTCAGCTCCGAGTAGCTGGGATTACAGGATCCACCACTAGTCTCGGCTAATAA 122874
QY 366 TGTITGTTTAAATTT---TTGAGGCCAGGTGCAAGTGGCCCATATCTGTGATCCAGCAC 422
Db 122975 AGTTGTTTTTAAAGTTGCTTCACGCTGGGTGCAGTGGCTCATGCTGTAAACCCAGCAC 123034
QY 423 TTTGGAGACCAAGCGAGCGCGATTTACTTGAAGTTCAGAGTTTCAAGACCGACCGAGCCAA 482
Db 123035 TTTGGAGCGCGAGCGAGCGAGATCACTGAGGTTCAGAGTTTCAGAGTACCTCTGGCCAA 123094
QY 483 CATGTAAACCATGTC 499
Db 123095 CATGCAAAACCCCATC 123111

Search completed: October 24, 2003, 21:05:57

Cob time : 1386.52 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 24, 2003, 15:39:20 Search time 125.999 Seconds
(without alignments)
10712.129 Million cell updates/sec

Title: US-09-830-902-1_COPY_1_500

Perfect score: 500

Sequence: 1 taactgactctgtgtgtttt.....aacatgtaaaacacatctg 500

Scoring table: IDENTITY_NUC

Gapop 10.0, Gapext 1.0

Searched: 2552756 seqs, 134979017 residues

Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database: N_Geneseq_19Jun03.*

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2: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARY

Result No.	Score	Query Match	Length	ID	Description
1	500	100.0	110000	22	AAF8480C Nucleoside sequence
2	175.6	35.1	57273	24	ABX22784 Human high bone ma
3	175.6	35.1	66933	25	ABA82625 Human HBV gene reg
4	175.6	35.1	66933	25	ACC45366 Human HBV gene fra
5	170.6	34.1	123722	24	ABQ98117 Human osteoblast d
6	169.2	33.8	33147	22	AAK57252 Human immune/haema
7	168.6	33.7	314	22	AAK33537 Human immune/haema
8	167.8	33.6	51469	22	AAK59322 Human immune/haema

C 9	167.8	33.6	51469	22	AAK70270 Human immune/haema
C 10	167.8	33.6	51469	22	AAK78813 Human immune/haema
C 11	167.2	33.4	2480	22	AAC85484 Human neurotrophin
C 12	166.6	33.3	249999	25	ABZ80229 Human tramodrin ge
C 13	165.8	33.2	32152	22	AAK39621 Genomic sequence #
C 14	165.8	33.2	32152	22	AAK39622 Human digestive sy
C 15	165.8	33.2	32152	22	AAK39623 Human digestive sy
C 16	165.8	33.2	32152	22	AAK91534 Human colorectal c
C 17	165.8	33.2	32152	22	AAK57799 Genomic DNA #172 e
C 18	165.8	33.2	32152	24	ABX59968 Genomic sequence #
C 19	165.8	33.2	32204	22	AAK39620 Human digestive sy
C 20	165.8	33.2	32204	22	AAK39621 Human digestive sy
C 21	165.8	33.2	32204	22	AAK15333 Human colorectal c
C 22	165.8	33.2	32204	24	ABX59967 Genomic DNA #171 e
C 23	165.8	33.2	36031	22	AAK66362 Human immune/haema
C 24	163.6	32.7	72349	24	ABX22783 Human high bone ma
C 25	163.6	32.7	72349	22	ABA82623 Human HBV gene reg
C 26	163.6	32.7	72349	25	ACC45364 Human HBV gene fra
C 27	163	32.6	9163	24	ABL45808 Human endothelia
C 28	162.4	32.5	14525	22	AAK70511 Human immune/haema
C 29	162.2	32.4	13024	22	AAK67339 Human reproductive
C 30	162.2	32.4	13024	23	ABL97644 Human testicular a
C 31	162	32.4	4447	22	AAK75249 Human immune/haema
C 32	160.8	32.2	1331	22	AAK66937 Human immune/haema
C 33	160.4	32.1	3459	22	AAK63901 Human digestive sy
C 34	160.4	32.1	3459	22	AAK19336 Human liver associ
C 35	160.4	32.1	1459	24	ABN90291 Human liver associ
C 36	160	32.0	84607	20	AAK59547 Human PACAP genom
C 37	157.8	31.6	8418	22	AAK41906 Genomic sequence #
C 38	157.6	31.5	1328	22	AAK66936 Human immune/haema
C 39	157	31.4	318	22	AAK62834 Human immune/haema
C 40	156.8	31.4	24606	22	AAK78367 Human immune/haema
C 41	156.8	31.4	24606	22	AAK94723 Human immune/haema
C 42	156.8	31.4	31051	22	AAK73223 Human immune/haema
C 43	156.2	31.2	1043	20	AAZ24862 Human secreted pro
C 44	155	31.0	10012	24	ABX55889 Human smalt induc
C 45	155	31.0	11096	22	ABA18278 Human nervous syst

ALIGNMENTS

RESULT 1
AAF8480C
10 AAF8480C standard; DNA; 110000 BP.
XX AAF8480C;
XX
XX 09-JUL-2001 (first entry)
XX Nucleotide sequence of the human SPG4 gene.
DE
DE Human; SPG4 gene; spastin; PSP-AD; gene therapy; s
XX autosomal dominant familial spastic paraplegia; s
XX Homo sapiens.

XX Key location/Qualifiers
FT CDS 9932..10209
FT exon /tag= a
FT /note= "contains introns"
FT /tag= b
FT /number= "1"
FT /tag= c
FT /number= "1"
FT /tag= d
FT /number= "2"
FT /tag= e
FT /number= "2"

RESULT 7
AAK83537/c
ID AAK83537 standard; DNA; 314 BP.
XX
XX AAK83537;
AC AAK83537;
CT 07-NOV-2001 (first entry)
XX Human immune/haematopoietic antigen germline sequence SEQ ID NO:38345.
DE Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
KW cytototoxic; gene therapy; vaccine; metastasis; ds.
XX Homo sapiens.
OS WC200157182-A2.
PN 09-AUG-2001.
XX 17-JAN-2001; 2001WO-US021354.
XX 31-JAN-2000; 2000US-0179065.
PR 24-FEB-2000; 2000US-0180628.
PR 24-FEB-2000; 2000US-0184564.
PR 22-MAR-2000; 2000US-0186350.
PR 16-MAR-2000; 2000US-0189374.
PR 17-MAR-2000; 2000US-0190376.
PR 18-APR-2000; 2000US-0198123.
PR 19-MAY-2000; 2000US-0205515.
PR 07-JUN-2000; 2000US-0209467.
PR 28-JUN-2000; 2000US-0214886.
PR 33-JUN-2000; 2000US-0215135.
PR 07-JUL-2000; 2000US-0216847.
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PR 01-DEC-2000; 200CUS-0250391.
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PR 08-DEC-2000; 200CUS-0251999.
PR 08-DEC-2000; 200CUS-0251999.
PR 11-DEC-2000; 200CUS-0254097.
PR 05-JAN-2001; 200CUS-0259678.
XX
XX (HUMAN): HUMAN GENOME SCI INC.
XX
XX Rosen CA, Barash SC, Ruben SM;
XX WPI; 2001-483426/52.
XX
XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
XX useful for preventing, diagnosing and/or treating cancers and
XX metastasis.
XX
XX Disclosure: SEQ ID NO 38349; 3071pp + Sequence Listing; English.
XX
XX AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
XX amino acid sequences given in AAK82170 to AAK91921. (I) have cytostatic
XX activity, and can be used in gene therapy and vaccine production. (I)
XX proteins and polynucleotides may be used in the prevention, diagnosis and
XX treatment of diseases associated with inappropriate (I) expression. For
XX example, they may be used to treat disorders associated with decreased
XX expression by rectifying mutations or deletions in a patient's genome
XX that affect the activity of (I) by expressing inactive proteins or to
XX supplement the patient's own production of (I). Additionally, (I)
XX polynucleotides may be used to produce the secreted (I), by inserting
XX the nucleic acids into a host cell and culturing the cell to express the
XX protein. (I) proteins and polynucleotides may be used to prevent,
XX diagnose and treat immune/haematopoietic-related diseases, especially
XX cancers and cancer metastases of haematopoietic-derived cells. AAK64703
XX to AAK87694 represent human immune/haematopoietic antigen genomic
XX sequences from the present invention. AAK54942 to AAK54950 and AAK54965
XX represent sequences used in the exemplification of the present invention.
XX
XX Sequence 314 BP; 86 A; 74 C; 87 G; 67 T; 0 other;
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XX Query Match: 32.7%; Score 169.6; DE 27; Length 314;
XX Best Local Similarity 74.8%; Fred. No. 4.6e-24;
XX Matches 225; Conservative 6; Mismatches 34; Indels 2; Gaps 17
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XX
XX 259 GATCTCAGCTCAGTGCAGCTCCACCTTCGGTTCGAAGCAATCCCTCCGCTCAATCT 318
XX ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
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XX ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX
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XX
XX 439 AGCCGATTAATTAGGTACGAGGTTCAGACACGACCGCCCAACATGGTAAACCATGT 498
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XX 3 C 3
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ID AAK69122 standard; DNA: 51469 BP.
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XX 06-NOV-2001 (first entry)
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XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:24134.
XX
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX cytostatic; gene therapy; vaccine; metastasis; ds.
XX
XX Homo sapiens.
XX
XX WC200157182-A2.
XX
XX 09-AUG-2001.
XX
XX 17-JAN-2001; 2001WC-US01354.
XX
XX 31-JAN-2003; 200CUS-C179065.
XX
XX 04-FEB-2003; 200CUS-0180628.
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XX 24-FEB-2003; 200CUS-0184664.
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XX 19-MAY-2003; 200CUS-C205515.
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XX 30-JUN-2003; 200CUS-C215135.
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XX 07-JUL-2003; 200CUS-C216647.
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XX 08-SEP-2003; 200CUS-0232080.
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XX 08-SEP-2003; 200CUS-0232081.
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ID AAK70270 standard; DNA; 51463 BP.
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AC AAK70270;
XX
DT 06-NOV-2003 (first entry)
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DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:25082.
XX
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX Cystostatic; gene therapy; vaccine; metastasis; GR.
XX
OS Homo sapiens.
XX
FN WO200157182-A2.
XX
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PF 17-JAN-2001; 2001WO-US01354.
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PR 01-DEC-2000; 2000US-0250391.
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PR 09-DEC-2000; 2000US-0251949.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254037.
PR 05-JAN-2001; 2001US-0259678.
XX (HUMA-): HUMAN GENOME SC: INC.
XX
XX Rosen CA, Barash SC, Ruben SM;
XX WPI; 2001-483426/52.
XX
XX Nucleic acids encoding human immune/haematopoietic antigen polypeptides,
XX useful for preventing, diagnosing and/or treating cancers and
XX metastasis.
XX
XX Disclosure: SEQ ID NO 25092; 3071pp + Sequence Listing: English.
XX
XX AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
XX amino acid sequences given in AAK82170 to AAK91922. They have cytostatic
XX activity, and can be used in gene therapy and vaccine production. (I)
XX proteins and polynucleotides may be used in the prevention, diagnosis and
XX treatment of diseases associated with inappropriate (I) expression. For
XX example, they may be used to treat disorders associated with decreased
XX expression by rectifying mutations or deletions in a patient's genome
XX that affect the activity of (I) by expressing inactive proteins or to
XX supplement the patient's own production of (I). Additionally, (I)
XX polynucleotides may be used to produce the secreted (I), by inserting
XX the nucleic acids into a host cell and culturing the cell to express the
XX protein. (I) proteins and polynucleotides may be used to prevent
XX cancers and treat immune/haematopoietic-related diseases, especially
XX cancers and cancer metastases of haematopoietic-derived cells. AAK64703
XX to AAK87694 represent human immune/haematopoietic antigen genomic
XX sequences from the present invention. AAK54942 to AAK54950 and AAK82169
XX represent sequences used in the exemplification of the present invention.
XX
XX Sequence 51469 BP; 13004 A; 13589 C; 13305 G; 1257 T; 0 other;
XX
XX Query Match 33.6%; Score 167.8; DB 22; Length 51469;
XX Best Local Similarity 73.6%; Pred. No. 1e-25; 77; Indels 5; Gaps 1;
XX Matches 229; Conservative 0; Mismatches 77;
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QY 194 TTTTATTTTATTTTCTAGATGGAGTCTTCTGTGGCCAGGCTGGAAATGCAAT 253
DB 16467 TTTTCTTTTCTTTTCTAGACAGAGTCTTCTGTGTGGCCAGGCTGGAGTGTAGT 16405
QY 254 GGCATGATCTCAGCTCATGTCAATCCACCTCCCTTGGGTTCAAGCAATCTCCCGGTCTCA 313
DB 16407 GGTGTGATCATAGCTCAGTACAGTCTCAAACTCTAGCTGAGAGATCTCTCCGGCTCA 16348
QY 314 ATCTCCAGTAGCTGGGACACAGAGATAGCTGCCACCTGGCTGGTTTCTTGT 373
DB 16347 GCATCCCAAGTAGCTGGGACTACTGTTATGCACACCTGGCTGGCTTATATAGCTCTC 16268
QY 374 TTTAA-----ATTTTGGCCAGGTGTCAGTGGCCCATATCTGTATCCAGCATTTTGGG 428
DB 16287 CTTAAGAATCTTCACTGGCTGGTGGTGGCTGATGCTGATCCAGCATTTGGG 16228
QY 429 AGACCAAGCAGCGCCGATTTACTTTAGGTTCAGGAGTCAAGACACCGCAGGCGACATGTT 488
DB 16227 AGCCCAAGCAGCAGATTTACCTGAGGTACAGGATTTGAGACACCGCTGACCAAGTGGT 16168
QY 489 AAAACCATGTC 499
DB 16167 GAACCCCTATC 16157
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RESULT 10
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XX 07-NOV-2001 (first entry);
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XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:13625.
XX
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX cytostatic; gene therapy; vaccine; metastasis; ds.
XX
XX Homo sapiens.
XX
XX NC00157182-A2.
XX
XX 09-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US01354.
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XX 31-JAN-2000; 2000US-0179365.
XX 04-FEB-2000; 2000US-0186628.
XX 24-FEB-2000; 2000US-0184664.
XX 02-MAR-2000; 2000US-0186350.
XX 16-MAR-2000; 2000US-0183874.
XX 17-MAR-2000; 2000US-0190076.
XX 18-APR-2000; 2000US-0191123.
XX 19-MAY-2000; 2000US-0205515.
XX 07-JUN-2000; 2000US-0209467.
XX 28-JUN-2000; 2000US-0214886.
XX 30-JUN-2000; 2000US-0215135.
XX 07-JUL-2000; 2000US-0215647.
XX 07-JUL-2000; 2000US-0216880.
XX 11-JUL-2000; 2000US-0217487.
XX 11-JUL-2000; 2000US-0217496.
XX 14-JUL-2000; 2000US-0218290.
XX 26-JUL-2000; 2000US-0220963.
XX 26-JUL-2000; 2000US-0220964.
XX 24-AUG-2000; 2000US-0224518.
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XX 14-AUG-2000; 2000US-0225213.
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XX 14-AUG-2000; 2000US-0225758.
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XX 22-AUG-2000; 2000US-0226681.
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XX 22-AUG-2000; 2000US-0227182.
XX 23-AUG-2000; 2000US-0227009.
XX 30-AUG-2000; 2000US-0228924.
XX 01-SEP-2000; 2000US-0229287.
XX 01-SEP-2000; 2000US-0229343.
XX 01-SEP-2000; 2000US-0229344.
XX 01-SEP-2000; 2000US-0229345.
XX 05-SEP-2000; 2000US-0229509.
XX 05-SEP-2000; 2000US-0229513.
XX 06-SEP-2000; 2000US-0231437.
XX 06-SEP-2000; 2000US-0231438.
XX 08-SEP-2000; 2000US-0231242.
XX 08-SEP-2000; 2000US-0231243.
XX 08-SEP-2000; 2000US-0231244.
XX 08-SEP-2000; 2000US-0231413.
XX 08-SEP-2000; 2000US-0231414.
XX 08-SEP-2000; 2000US-0231420.
XX 08-SEP-2000; 2000US-0232381.
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RESULT 11
AAC85484/c
ID AAC85484 standard; cDNA, 2480 BP.
XX
AC AAC85484;
XX
DT 16-MAY-2001 (first entry)
XX
DE Human neuropeptide FF promoter.
XX
KW Promoter; murine; neuropeptide FF; NPFF; brain; spinal cord; alacrima;
KW inflammation; transcription factor; NPFFa/b; hormonal dysfunction;
KW nuclear factor of activated T-cells; NFAT; heat shock factor 1; ACTH;
KW HSF1; Allgrove syndrome; triple-A syndrome; sensory impairment;
KW adrenocorticotrophic hormone; resistant adrenal insufficiency;
KW achalasia; hypoglycaemia; autonomic neuropathy; autonomic function;
KW gene therapy; ds.
XX
OS Homo sapiens.
XX
FH Key location/qualifiers
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FT protein_bind /bound_moiety= "AP1"
FT protein_bind complement (2122..2128)
FT protein_bind /*tag= b
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FT protein_bind 2155..2171 /*tag= c
FT protein_bind /bound_moiety= "QMYB"
FT protein_bind 2206..2216 /*tag= d
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FT /*tag= h
XX
EPIC74619-A2.
XX
PD 07-FEB-2001.
XX
XX 28-JUL-2000; 2000EP-0660130.
XX
XX 03-AUG-1999; 99US-0365755.
XX 27-MAR-2000; 2000US-0534639.
XX
PA (PANU/; PANULA P A C.
PA (BRAN/; BRANDT A.
PA (WEST/; WESTERLUND J.
XX
P1 Panula PAJ, Brandt A, Westerlund J;
XX
XX WPI; 2001-171047/18.
XX
XX New promoter for neuropeptide FF (NPFF), useful for treating and
PT screening genetic diseases associated with NPFF gene promoter such as
PT deficient regulation of autonomic function and pain conditions -
XX
PS Claim 2; Fig 4; 55pp; English.
XX
XX This sequence represents the promoter region for the human neuro-
CC peptide FF (NPFF) gene. The NPFF gene is expressed in specific regions
CC of the brain and in the spinal cord and is induced upon inflammatory
CC stimulus. Binding sites were found for the inflammation related

transcription factor, NFkappaB, and for the nuclear factor of
activated T-cells (NFAT). A binding site was also found for heat
shock factor 1 (HSF1) which is activated in cells at elevated
temperatures and other environmental stress conditions. The
AC-dinucleotide repeat is thought to add additional regulatory
effects. The human NPFF gene is located in the human chromosome
locus 12q13 which is known to be involved in Allgrove syndrome
(triple-A syndrome) which is characterised by a triad of adreno-
corticotrophic hormone (ACTH), resistant adrenal insufficiency,
achalasia and alacrima, hypoglycaemia and sensory impairment; and
autonomic neuropathy. The NPFF promoter region may be useful for
treating genetic diseases such as those associated with deficient
regulation of autonomic function, pain conditions or hormonal
dysfunction which are associated with the promoter area of the
NPFF gene whose expression is modulated through the regulatory sites.
It is also useful in the screening of the genetic diseases associated
with the promoter area of the NPFF gene by modulation of activation or
inhibition of NPFF gene expression through the regulation sites in the
promoter area, and is used in gene therapy and DNA analysis.
XX
XX Sequence 2480 BP; 638 A; 536 C; 673 G; 573 T; 0 other;
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Query Match 33.4%; Score 167.2; DB 22; Length 2480;
Best Local Similarity 75.6%; Pred. No. 11e-25;
Matches 239; Conservative 0; Mismatches 63; Indels 14; Gaps 2;
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DB 1885 AATTTAACTTTTCTTTTTCAGATGGAGTTTCCCTTTTTCGCCAGGCTGGAGTCCA 1826
QY 252 ATGGCATGATCTCACTCATCTGCAACCTCCAGCTTTGGGTTTCAGGCAATCTCTGGCT 311
DB 1825 GTGGCGGCTCTGGCTCACTGCAACCTCTGGCTCTGGGTTCAAGTATTCTCTGGCT 1768
QY 312 CAATCTTCAGTACTGGACTACAGCAATGAGCTGCGGACCTGGCTGGTTGGTTG 371
DB 1765 CAGCTCCCAAGTAGCTGGGACTACAGGCGTCAACACCATGCCAGGCTGATCTTTT 1706
QY 372 TTTTAA-----TTTTSAGGCGAGTGGAGTGGCGCATATCTGTATGCCA 418
DB 1705 TTTTAAATAAACCATTTTTTCGGGGTGGAGTGGCTGACACCTGTACCCCA 1646
QY 419 GCATTTGGGAGACCAAGCGAGGCGGATTAATTTAGGTACAGAGTTCAAGACCCAGG 478
DB 1645 GCATTTGGGAGACCAAGCGGCGGATCACTGAGGCGAGAGTCGAGACCGCTGG 1586
QY 479 CCAACATGGTAAACC 494
DB 1585 CCAACATGGTAAACC 1570
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XX
XX 02-JUN-2003 (first entry)
XX
DE Human tramdorin gene region genomic DNA SEQ ID NO:26.
XX
KW Neuroprotective; nootropic; cerebroprotective; analgesic; gene therapy;
KW central nervous system disorder; CNS disorder; multiple sclerosis;
KW nerve injury; neuropathic pain; stroke; trauma; non-CNS disorder; tramd;
KW tramdorin; human; chromosome 5; gene; ds.
XX
OS Homo sapiens.
XX
XX NC020307.6522-A2.
XX
XX 27-FEB-2003.
XX
XX 21-AUG-2002; 2002MC-US26637.
XX

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Without alignments:
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Title: US-09-830-902-1_COPY_1_500

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Scoring table: IDENTITY_NJC

Gapop 10.0 , Gapext 1.0

Searched: 569378 seqs, 220691566 residues

Total number of hits satisfying chosen parameters: 1139056

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 2%

Maximum Match 100%

Listing first 45 summaries

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- 5: /cdn2_6/prodata/2/ina/ACTUS_COV3.seq:*
- 6: /cdn2_6/prodata/2/ina/backfiles.seq:*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	167.2	33.4	2480	US-09-534-638-3	Sequence 3, Appl 1
C 2	141.2	25.2	18000	US-09-657-346A-17	Sequence 17, Appl 1
C 3	136.4	27.3	45716	US-08-965-048-5	Sequence 5, Appl 1
C 4	136.4	27.3	45989	US-08-965-048-6	Sequence 6, Appl 1
C 5	136.4	27.3	174493	US-09-804-47A-3	Sequence 3, Appl 1
C 6	133.6	26.7	2268	US-09-369-247-42	Sequence 42, Appl 1
C 7	133.6	26.7	75395	US-09-984-890-3	Sequence 3, Appl 1
C 8	132.6	26.5	319608	US-09-539-333D-1	Sequence 1, Appl 1
C 9	132.6	26.5	319608	US-09-679-409-1	Sequence 1, Appl 1
C 10	132.4	26.5	40000	US-09-780-049-18	Sequence 18, Appl 1
C 11	132	26.4	74962	US-09-685-853A-3	Sequence 3, Appl 1
C 12	132	26.4	90541	US-09-759-359A-3	Sequence 3, Appl 1
C 13	131.6	26.3	71310	US-09-056-105-31	Sequence 31, Appl 1
C 14	131.2	26.2	66804	US-09-74C-041-3	Sequence 3, Appl 1
C 15	131	26.2	1701	US-09-078-294-9	Sequence 9, Appl 1
C 16	130.8	26.2	55298	US-09-491-356C-1	Sequence 1, Appl 1
C 17	129.8	26.0	23187	US-09-499-522-1	Sequence 1, Appl 1
C 18	129.4	25.9	19650	US-09-819-989-3	Sequence 3, Appl 1
C 19	129.4	25.9	80246	US-09-078-294-4	Sequence 4, Appl 1
C 20	129.2	25.8	3885	US-08-688-145-1	Sequence 1, Appl 1
C 21	129.2	25.8	116592	US-09-818-512-3	Sequence 3, Appl 1
C 22	129	25.8	5375	US-08-757-223-7	Sequence 7, Appl 1
C 23	129	25.8	82595	US-09-078-294-3	Sequence 3, Appl 1
C 24	129	25.8	116592	US-09-818-512-3	Sequence 3, Appl 1
C 25	128.4	25.7	3002	US-09-539-333D-218	Sequence 218, Appl 1
C 26	128	25.6	863	US-08-943-731-171	Sequence 171, Appl 1
C 27	128	25.6	17606	US-08-943-731-4	Sequence 4, Appl 1

28	127.8	25.6	1754	4	US-09-620-312D-518	Sequence 518, Appl 1
29	127.8	25.6	4754	2	US-08-476-062A-52	Sequence 52, Appl 1
30	127.8	25.6	20201	4	US-09-734-674-3	Sequence 3, Appl 1
C 31	127.6	25.5	16063	4	US-09-801-052-3	Sequence 3, Appl 1
C 32	127	25.4	3463	4	US-09-304-615-44	Sequence 44, Appl 1
C 33	127	25.4	55288	4	US-09-491-356C-1	Sequence 1, Appl 1
C 34	126.8	25.4	62834	4	US-09-800-960-3	Sequence 3, Appl 1
C 35	126.4	25.3	8596	3	US-08-943-731-31	Sequence 31, Appl 1
C 36	126.4	25.3	16609	3	US-08-943-731-1	Sequence 1, Appl 1
C 37	126.4	25.3	99500	4	US-09-796-096-10	Sequence 10, Appl 1
C 38	126.2	25.2	148567	4	US-09-801-876B-3	Sequence 3, Appl 1
C 39	126	25.2	50000	4	US-09-146-053-4	Sequence 4, Appl 1
C 40	126	25.2	19638	4	US-09-539-333D-1	Sequence 1, Appl 1
C 41	126	25.2	19638	4	US-09-679-409-1	Sequence 1, Appl 1
C 42	125.6	25.1	55927	4	US-09-823-133A-3	Sequence 3, Appl 1
C 43	125.4	25.1	1000	3	US-09-018-584A-23	Sequence 23, Appl 1
C 44	125.4	25.1	2059	3	US-08-938-669A-5	Sequence 5, Appl 1
C 45	125.4	25.1	2059	4	US-09-306-828-5	Sequence 5, Appl 1

ALIGNMENTS

Result :
US-09-534-638-3/c
Sequence 3, Application US/09534638
Recent No. 6320038
GENERAL INFORMATION:
APPLICANT: Panula, Pertti A.J.
APPLICANT: Brandt, Anika
APPLICANT: Westerlund, Johanna
TITLE OF INVENTION: Promoter for Neurotrophin FF Promoter and use thereof
TITLE OF INVENTION: for therapy and diagnosis
FILE REFERENCE: 2530-104
CURRENT APPLICATION NUMBER: US/09/534,638
CURRENT FILING DATE: 2000-03-27
EARLIER APPLICATION NUMBER: 09/365755
PAPER FILING DATE: 1999-08-03
NUMBER OF SEQ ID NOS: 22
SOFTWARE: Patent In Ver. 2.1
SEQ ID NO 3
LENGTH: 2480
TYPE: DNA
ORGANISM: Homo sapiens
US-09-534-638-3

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Gaps	2						
Oy	193	ATTTATTTTATTTTATTTTCTAGTGGAGCTTGCTC-TGTTGCCAGGCTGGAATGCA	251				
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Oy	252	ATGGCATGATCTCAGCTCACTGCAACCTTCACCTTTTGGGTTTCAAGCAATCTCTCGCT	311				
De	1825	GTGGCGGAGTCTGGGCTCACTGCAACCTCTGGTCTCTGGTTCAGTGATTCCTCGCT	1766				
Oy	312	CAATCTTCCAGTAGCTGGGACTACAGGAATGAGCTGCCGACCTGGCTGGTTGTTTG	371				
De	1765	CAGCTCCCAAGTAGCTGGGACTACAGGCTCAACCACTGCCCCAGCTGTATCTTTT	1706				
Oy	372	TTTTTAA-----TTTTGAGGCGAGTGCAGTGGCCCATATCTGTATCCCA	418				
De	1705	TTTTTAA-----TTTTGAGGCGAGTGCAGTGGCCCATATCTGTATCCCA	1646				
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De	1645	GCACTTTGGAGACCAAGGAGCGCCGATTTACTTGGGTTCAGGAGTTCAGACACGAGCAG	1566				
Oy	479	CCACATGCTAAACCC	494				
De	1585	CCACATGCTAAACCC	1570				


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RESULT 2
US-09-657-346A-17
; Sequence 17, Application US/09657346A
; Patent No. 6503754
; GENERAL INFORMATION:
; APPLICANT: Hong Zhang
; APPLICANT: Jacqueline Wyatt
; TITLE OF INVENTION: ANTISENSE MODULATION OF BDN INTERACTING DOMAIN DEATH AGONIST
; TITLE OF INVENTION: EXPRESSION
; FILE REFERENCE: RTS-0135
; CURRENT APPLICATION NUMBER: US/09/657,346A
; CURRENT FILING DATE: 2000-09-07
; NUMBER OF SEQ ID NOS: 174
; SEQ ID NO 17
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; TYPE: DNA
; ORGANISM: Homo sapiens
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; NAME/KEY: CDS
; LOCATION: (2144)...(2155)
; NAME/KEY: CDS
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US-09-657-346A-17

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QY 203 ATTATTTTCTAGATGGAGCTTGTCTGTCTCCAGGCTGGAATGCAATGGCATGATC 262
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QY 263 TCGCTCTAGTCAAGCTTCCAGCTTGGGTTCAGCAATCTCTCTCCCAATCTTCCAA 322
DB 11415 ACAGCTCACGGACCTTGAATCTTGGCTCAAGAGTCTCTCCCACTCCAGCTCTCT 11474
QY 323 GTAGCTGGGACTACGAAATAGCTCCGACCTGCTGCTGCTGCTGCTGCTGCTGCT 382
DB 11475 GTAGCTGGGACTACGAAATAGCTCCGACCTGCTGCTGCTGCTGCTGCTGCTGCT 11520
QY 383 TGAGGCGAGCTGGAGTGGGCTATATTTGTATCCAGGCTTGGAGACCAAGGACAGC 442
DB 11531 AGAGGCGAGGACAGTGGCTCACACTGTATCCAGGCTTGGAGACCAAGGACAGC 11590
QY 443 CGATTACTTGGTCAAGATTCAGACACGCGAGGCGGACATGCTGTAAGAC 494
DB 11591 GGAATCAG--AAGGTGAGGATTCGAGACAGCGCTGACCAACATGGTGAAGCC 11640

RESULT 3
US-08-965-048-5
; Sequence 5, Application US/08965048
; Patent No. 6323244
; GENERAL INFORMATION:
; APPLICANT: Chen, Hong
; APPLICANT: Freimer, Nelson
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DIAGNOSIS AND
; TITLE OF INVENTION: TREATMENT OF NEUROPSYCHIATRIC DISORDERS
; FILE REFERENCE: 7853-093
; CURRENT APPLICATION NUMBER: US/08/965,048
; CURRENT FILING DATE: 1997-11-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 5
; LENGTH: 45989
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-08-965-048-6

Query Match      27.3%; Score 136.4; DB 4; Length 45989;
Best Local Similarity 71.2%; Pred. No. 7.2e-23;
Matches 195; Conservative 0; Mismatches 76; Indels 3; Gaps 1;
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; TYPE: DNA
; ORGANISM: Homo sapiens
US-08-965-048-5

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Best Local Similarity 71.2%; Pred. No. 7.2e-23;
Matches 195; Conservative 0; Mismatches 76; Indels 3; Gaps 1;

QY 106 CATTTTCACCTGAAGTAAACAATCTCCATAAATCTCCATAAATCTCAAACTGCTGCTGGGA 165
DB 34614 CAATTGAGGTTAATAGACATGCTCTCTATGCTTTGTAATACAGAAATCTGACACCAT 34673
QY 166 GTTTTGGTTTGTCTTTTGTCTTTTATTTTATTTTATTTTATTTTCT---AGATGGAG 222
DB 34674 TTTTCTTTTGTCTTTTGTCTTTTATTTTATTTTATTTTATTTTCTTTTGTGATAGAG 34733
QY 223 TCTTGTCTGTGTTGCCAGGCTGGAATGCAATGGCATGATCTGAGTCACTGCACTGCACTCCA 282
DB 34734 TCTTACTCTGTGTTGCCAGGCTGGAATGCAATGGCATGATCTGAGTCACTGCACTCCA 34793
QY 283 CCTTCGGGTTCAAGCAATCTCTCTGCTCAATCTTCCAACTAGCTGAGTGGAGTACAGGAAT 342
DB 34794 CTTCCGAGTTCAAGCAATCTCTCTGCTCAAGCTCTGAGTGGAGTGGAGTACAGATGT 34853
QY 343 GAGCTGGCGACCTGGCTGGTGTGTTGTTT 376
DB 34854 GTGCCACCATGCTGGCTTTTCTTTTCTTTT 34897

RESULT 4
US-08-965-048-6
; Sequence 6, Application US/08965048
; Patent No. 6323244
; GENERAL INFORMATION:
; APPLICANT: Chen, Hong
; APPLICANT: Freimer, Nelson
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DIAGNOSIS AND
; TITLE OF INVENTION: TREATMENT OF NEUROPSYCHIATRIC DISORDERS
; FILE REFERENCE: 7853-093
; CURRENT APPLICATION NUMBER: US/08/965,048
; CURRENT FILING DATE: 1997-11-05
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 6
; LENGTH: 45989
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-08-965-048-6

Query Match      27.3%; Score 136.4; DB 4; Length 45989;
Best Local Similarity 71.2%; Pred. No. 7.2e-23;
Matches 195; Conservative 0; Mismatches 76; Indels 3; Gaps 1;

QY 106 CATTTTCACCTGAAGTAAACAATCTCCATAAATCTCCATAAATCTCAAACTGCTGCTGGGA 165
DB 34728 CAATTGAGGTTAATAGACATGCTCTCTATGCTTTGTAATACAGAAATCTGACACCAT 34787
QY 166 GTTTTGGTTTGTCTTTTGTCTTTTATTTTATTTTATTTTATTTTCT---AGATGGAG 222
DB 34786 TTTTCTTTTGTCTTTTGTCTTTTATTTTATTTTATTTTATTTTCTTTTGTGATAGAG 34847
QY 223 TCTTGTCTGTGTTGCCAGGCTGGAATGCAATGGCATGATCTGAGTCACTGCACTGCACTCCA 282
DB 34848 TCTTACTCTGTGTTGCCAGGCTGGAATGCAATGGCATGATCTGAGTCACTGCACTCCA 34907
QY 283 CCTTCGGGTTCAAGCAATCTCTCTGCTCAATCTTCCAACTAGCTGAGTGGAGTACAGGAAT 342
DB 34908 CTTCCGAGTTCAAGCAATCTCTCTGCTCAAGCTCTGAGTGGAGTGGAGTACAGATGT 34967
QY 343 GAGCTGGCGACCTGGCTGGTGTGTTGTTT 376
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/ EARLIER FILING DATE: 1988-02-09
/ NUMBER OF SEQ ID NOS: 172
/ SOFTWARE: Patentin Ver. 2.0
/ SEQ ID NO 42
/ LENGTH: 1269
/ TYPE: DNA
/ ORGANISM: Homo sapiens
/ FEATURES:
/ NAME/KEY: SITE
/ LOCATION: (2)
/ OTHER INFORMATION: n equals a.t.g. or c
/ FEATURE:
/ NAME/KEY: SITE
/ LOCATION: (15)
/ OTHER INFORMATION: n equals a.t.g. or c
/ FEATURE:
/ NAME/KEY: SITE
/ LOCATION: (23)
/ OTHER INFORMATION: n equals a.t.g. or c
/ FEATURE:
/ NAME/KEY: SITE
/ LOCATION: (368)
/ OTHER INFORMATION: n equals a.t.g. or c
/ FEATURE:
/ NAME/KEY: SITE
/ LOCATION: (479)
/ OTHER INFORMATION: n equals a.r.g. or c
/ US-09-369-247 42

Query Match          26.7%   Score 135.6; DS 4; Length 1268;
Best Local Similarity 79.4%; Pred. No. 1.ee-22;
Matches 160; Conservative
Gaps 0; Indels 0; Mismatches 44;

QY      173  TTTTGGTTTTTGTTGTTTAATTTATTTATTATTTCTAGATGGAGTTTCGGCG 232
DB      1260  TTTTGTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTGACGAGTTTCTGTG 1261

QY      233  TTGCCAGGCTGGAAATGCATATCCAGTCACTGCAACTCAGCTTTCGGT 292
DB      1260  TGCCGAGGTGGAGTGCACTGGCGGAATCTGGGTCACTGCAAGCTCCGGCTCCGGGC 1261

QY      293  TCAGCAATCTCTCGGCTCAATCTTCAGTAGTCTGGAGTACAGSAAATGAGTTCGGC 362
DB      1140  TCAGCGCATCTCTCGGCTCAGCTCCGAGTAGTGGAGTACAGCGCGCGCTACAC 1681

QY      363  ACCCGGGCTGTTTGTGTGTTTT 376
DB      1080  GCCCGGTAAATTCTTGATTTTT 1057

RESULT 7
US-09-984-890-3/c
/ Sequence 3, Application US/09984890
/ Patent No. 6492156
/ GENERAL INFORMATION:
/ APPLICANT: YAN, Chunhua et al.
/ TITLE OF INVENTION: ISOLATED HUMAN KINASE PROTEINS, NUCLEIC
/ TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN KINASE PROTEINS, AND USES
/ TITLE OF INVENTION: THEREOF
/ FILE REFERENCE: CLC01306
/ CURRENT APPLICATION NUMBER: US/09/964,890
/ CURRENT FILING DATE: 2001-10-31
/ NUMBER OF SEQ ID NOS: 4
/ SOFTWARE: FastSeq for Windows Version 4.0
/ SEQ ID NO 3
/ LENGTH: 75395
/ TYPE: DNA
/ ORGANISM: Homo sapiens
/ FEATURES:
/ NAME/KEY: misc_feature
/ LOCATION: (1)...(75395)
/ OTHER INFORMATION: n = A,T,C or G
/ US-09-984-890-3

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Query Match      26.7%   Score 133.6; DB 4; Length 75325;
Sest Local Similarity 72.6%; Pred. No. 3,5e+22;
Matches 215; Conservative 0; Mismatches 74; Indels 7; Gaps 3;

QY 204 TTTTATTTTCAGATGGATGCTGCTCTGTCGCGAGCGGTGAAATGCAATGGCATGATCT 263
DB 10020 TATTATCTTGATGGGTCTCACTCTCTGCCCAGGCTGGAAATGCAATGGCATGATCA 9961

QY 264 CAGCTCACTCAACCTCCAGCTTTCGGGTCAAGCAATCTCTGCTGCTCAATCTTCGAG 323
DB 9960 CAGCTCACTCAACCTCCAGCTTTCGGGTCAAGCAATCTCTGCTGCTCAATCTTCGAG 9961

QY 324 TAGCTGGGACTACAGAAATGAGCTGCGGCACTGCGCTGCTGTTTGTGTTTATATTIT 383
DB 9900 TAGGTGGGAATACACCATGCTGTACACACACACACACACACACACACACACACACAC 9965

QY 384 GAGGCGAGGTGGAGTGGAGTGGAGTGGAGTGGAGTGGAGTGGAGTGGAGTGGAGTGG 443
DB 9845 TGGCTGGGCGGGGGTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGGCTGG 9785

QY 444 GATTACTTGAAGTCAAGAGTTCAAGACCGAGCGGCGGCAATGTTAAACCATGTC 499
DB 9785 GATCACACAGGTCAAGAGTCAAGACCGAGCGGCGGCAATGTTAAACCATGTC 9732

RESULT 8
US-09-539-333D-1/2
; Sequence 1, Application US/09549333D
; Patent No. 6476208
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; APPLICANT: Bougueleret, Lydie
; APPLICANT: Bihain, Bernard
; APPLICANT: Essioux, Laurent
; TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENES, PROTEINS AND BIALLIC MARKERS
; FILE REFERENCE: GENSET.547AUS
; CURRENT APPLICATION NUMBER: US/09/539,333D
; PRIOR FILING DATE: 2000-03-30
; PRIOR APPLICATION NUMBER: US 60/126,903
; PRIOR FILING DATE: 1999-03-30
; PRIOR APPLICATION NUMBER: US 60/131,971
; PRIOR FILING DATE: 1999-04-30
; PRIOR APPLICATION NUMBER: US 60/132,065
; PRIOR FILING DATE: 1999-04-30
; PRIOR APPLICATION NUMBER: US 60/143,928
; PRIOR FILING DATE: 1999-07-14
; PRIOR APPLICATION NUMBER: US 60/145,915
; PRIOR FILING DATE: 1999-07-27
; PRIOR APPLICATION NUMBER: US 60/146,453
; PRIOR FILING DATE: 1999-07-29
; PRIOR APPLICATION NUMBER: US 60/146,452
; PRIOR FILING DATE: 1999-07-29
; PRIOR APPLICATION NUMBER: US 60/142,288
; PRIOR FILING DATE: 1999-10-28
; PRIOR APPLICATION NUMBER: US 60/146,364
; PRIOR FILING DATE: 1999-10-12
; NUMBER OF SEQ ID NOS: 23;
; SOFTWARE: Patent.pm
; SEQ ID NO 1
; LENGTH: 319608
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: 31..1107
; OTHER INFORMATION: 5' regulatory region g35018 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 1108..1289
; OTHER INFORMATION: exon A g35018 gene

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; FEATURE:
; NAME/KEY: exon
; LOCATION: 14877..14920
; OTHER INFORMATION: exon B g35018 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 18778..18862
; OTHER INFORMATION: exon Bb1s g35018 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 25593..25740
; OTHER INFORMATION: exon C g35018 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 29389..29502
; OTHER INFORMATION: exon D g35018 gene
; FEATURE:
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; OTHER INFORMATION: exon E g35018 gene
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; LOCATION: 64666..64812
; OTHER INFORMATION: exon F g35018 gene
; FEATURE:
; NAME/KEY: exon
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; OTHER INFORMATION: 3' regulatory region g35018 gene
; FEATURE:
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; LOCATION: 94124..94964
; OTHER INFORMATION: exon g35017
; FEATURE:
; NAME/KEY: exon
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; OTHER INFORMATION: exon S g35030 gene
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; OTHER INFORMATION: exon T g35030 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 215702..215746
; OTHER INFORMATION: exon U g35030 gene
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; NAME/KEY: exon
; LOCATION: 216836..216915
; OTHER INFORMATION: exon V g35030 gene
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: 213818..215818
; OTHER INFORMATION: 3' regulatory region g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 215829..215941
; OTHER INFORMATION: exon R complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 215819..215975
; OTHER INFORMATION: exon Bb1s complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 216661..216952
; OTHER INFORMATION: exon Qb1s complement g34872 gene
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; NAME/KEY: exon
; LOCATION: 216661..217061
; OTHER INFORMATION: exon Q complement g34872 gene
; FEATURE:

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; NAME/KEY: exon
; LOCATION: 217027..217061
; OTHER INFORMATION: exon Q1 complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 229647..229742
; OTHER INFORMATION: exon X complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 230408..230721
; OTHER INFORMATION: exon P complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 231272..231412
; OTHER INFORMATION: exon Obis complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 231787..231890
; OTHER INFORMATION: exon O2 complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 231870..231879
; OTHER INFORMATION: exon O1 complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 234174..234321
; OTHER INFORMATION: exon O complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 237406..237428
; OTHER INFORMATION: exon Nbis complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 239719..239807
; OTHER INFORMATION: exon N2 complement g34872 gene
; FEATURE:
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; LOCATION: 239719..239853
; OTHER INFORMATION: exon N complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 240528..240569
; OTHER INFORMATION: exon M117 complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 240528..240596
; OTHER INFORMATION: exon M1000 complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 240528..240644
; OTHER INFORMATION: exon X82 complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 240528..240824
; OTHER INFORMATION: exon X862 complement g34872 gene
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; LOCATION: 240528..240994
; OTHER INFORMATION: exon M692 complement g34872 gene
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; NAME/KEY: exon
; LOCATION: 240528..241685
; OTHER INFORMATION: exon M1 complement g34872 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 240900..240993
; OTHER INFORMATION: exon M51 complement g34872 gene
; FEATURE:
; NAME/KEY: misc_feature

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; LOCATION: 241686..243685
; OTHER INFORMATION: 5'regulatory region: g34872 gene
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 290652..292652
; OTHER INFORMATION: 3'regulatory region g34665 gene
; FEATURE:
; NAME/KEY: exon
; LOCATION: 292653..292841
; Query Match: 26.5%; Score 132.6; DB 4; Length 319608;
; Best Local Similarity 76.8%; Pred. No. 8.1e-22;
; Matches 162; Conservative 0; Mismatches 43; Indels 2; Gaps 0;
Qy 173 TTTTGTTTTTCTGTTTATTTTATTTTATTTTATTTTCTAGATGGAGTGTTCCTCTG 332
Db 16340 TTTTCGTTCTGTTGTTGTTTGTGTTTGTGTTTGTGTTTGTGTTGTTGTTGTTG 16281
Qy 233 TTGCCCAGGCTGGAATGCAATGCAATGATCTCAGCTCACTGCAACCTCCACCTTTGGGT 292
Db 16280 TTGCTCAGGCTGGAGTGTAGTSCGCGATCTCAGCTCACTGCAACCTTCGCTCCCGGT 16221
Qy 293 TCAAGCAATCTCTCCTCAATCTTCCAAAGTAGCTGGGACTACAGGAATGAGTCCGCC 352
Db 16220 TCAAGCAATCTCATGACTCAGCTCCCAAGTAGCTGGGATACAGGCGCATGCCACCA 16161
Qy 353 ACGTGGCTGGTTGTTGTTGTTGTTTAAATTT 383
Db 16160 GCGTGGCTAAATTTTCTGTTGTTT 16130
RESULT 9
US-09-679-409-1/C
; Sequence 1, Application US/09679409
; Patent No. 655316
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Vania
; APPLICANT: Grumakov, Ilya
; APPLICANT: Bengueler, Lydie
; APPLICANT: Essioux, Laurent
; TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENE, PROTEINS AND BIALLLELIC MARKER
; FILE REFERENCE: 53-US15,61P
; CURRENT APPLICATION NUMBER: US/09/679,409
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 09/539,333
; PRIOR FILING DATE: 2000-03-03
; PRIOR APPLICATION NUMBER: 09/416,384
; PRIOR FILING DATE: 1999-10-12
; PRIOR APPLICATION NUMBER: 60/168,088
; PRIOR FILING DATE: 1999-11-30
; NUMBER OF SEQ ID NOS: 134
; SOFTWARE: Patent.pm
; SEQ ID NO 1
; LENGTH: 319608
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 199122..201122
; OTHER INFORMATION: 5'regulatory region
; NAME/KEY: exon
; LOCATION: 201123..201234
; OTHER INFORMATION: exon S
; NAME/KEY: exon
; LOCATION: 201123..201569
; OTHER INFORMATION: exon S2
; NAME/KEY: exon
; LOCATION: 214676..214793
; OTHER INFORMATION: exon T
; NAME/KEY: exon
; LOCATION: 215702..215746
; OTHER INFORMATION: exon U

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1 : Sequence 1411, Application US/10121741
2 : Publication No. US20030143668A1
3 : GENERAL INFORMATION:
4 : APPLICANT: SUWA, MAKIKO
5 : APPLICANT: ASAI, KIYOSHI
6 : APPLICANT: AKIYAMA, YUTAKA
7 : APPLICANT: ABURATANI, HIROYUKI
8 : TITLE OF INVENTION: NOVEL G PROTEIN-COUPLED RECEPTORS
9 : F-FILE REFERENCE: C84335/C-152
10 : CURRENT APPLICATION NUMBER: US/010117.141
11 : CURRENT FILING DATE: 2002-12-18
12 : PRIOR APPLICATION NUMBER: JP 2001/246789
13 : PRIOR FILING DATE: 2001-06-18
14 : NUMBER OF SEQ ID NOS: 2430
15 : SOFTWARE: PatentIn Ver. 2.1.1
16 : SEQ ID NO 1611
17 : LENGTH: 39725
18 : TYPE: DNA
19 : ORGANISM: Homo sapiens
20 : FEATURE:
21 : NAME/KEY: source
22 : LOCATION: (1)..(39725)
23 : FEATURE:
24 : NAME/KEY: CDS
25 : LOCATION: (201)..(409)
26 : FEATURE:
27 : NAME/KEY: CDS
28 : LOCATION: (4795)..(12966)
29 : FEATURE:
30 : NAME/KEY: CDS
31 : LOCATION: (3861)..(4281)
32 : FEATURE:
33 : NAME/KEY: CDS
34 : LOCATION: (6128)..(6513)
35 : FEATURE:
36 : NAME/KEY: CDS
37 : LOCATION: (13450)..(13658)
38 : FEATURE:
39 : NAME/KEY: CDS
40 : LOCATION: (15063)..(15253)
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43 : LOCATION: (15939)..(16148)
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46 : LOCATION: (20042)..(20167)
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66 : NAME/KEY: CDS
67 : LOCATION: (28906)..(29142)
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69 : NAME/KEY: CDS
70 : LOCATION: (30237)..(30375)
71 : FEATURE:
72 : NAME/KEY: CDS
73 : LOCATION: (36366)..(36636)

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QY 240 GCGTGGATGCAATGGCATGATCTCAGCTCAGTGGCAACCTCCAGCTTTGGGCTTCAAGCA 239
DB 211140 GCGTGAAGTSCAGTGGCAATCAACAGATTCATCTGTGACCTCTCTTCCAGGCTCAAGCA 211081
QY 300 ATCTCTCTCTCCCTCAATCTTCCAGTAGCTGGGACTACAGGAATGAGCTGCCGCACTGGC 359
DB 211080 ATCTCTCAACCTCAGCTCTCTGAGTAGTGGGACTACAGGATTCACCATCATGAGCCAGC 211021
QY 360 CTGGTTTGTGTTTTTAA-----TTTGGAGCCAGTGGAGTGGGCCATATCTG 410
DB 211020 TAAATTTTGTATCTCTGTAGATATGGGTTTTGTGGCCGGTGGAGTGGCTCATGGCTG 210961
QY 411 TGAATCCAGCACTTTGGGAGAGCAAGGAGCGGATTAATCTCAGGTTCAGGACTTCAAGAC 470
DB 210960 TAAATCCAGCACTTTGGGAGGCAAGGTGGGAGATCTCAGGTTCAGGACTTCAAGAC 210901
QY 471 CAGGCGAGGCAACATGTATAAACCATGTC 499
DB 210900 CAGGCTGGCCAAACATGTGTAAACCCCATC 210872

RESULT 8
US-09-764-855-328
; Sequence 328, Application US/09764855
; Patent No. US20020115919A;
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: P1101
; CURRENT APPLICATION NUMBER: US/09/764,855
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult P1101 or file wrapper
; NUMBER OF SEQ ID NOS: 334
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 328
; LENGTH: 32152
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-855-328

Query Match 33.2%; Score 165.8; DB 10; Length 32152;
Best Local Similarity 74.6%; Pred. No. 4.6e-32;
Matches 244; Conservative 0; Mismatches 62; Indels 21; Gaps 2;

QY 192 TATTTATTTATTTATTTATTTCTAGATGGAGTCTGCTGTGGCCAGGCTGGAAATGCA 251
DB 3262 TGTTTTTTTTTTTTTTTTTTTTGGAGCGGAGTCTGCTGTGGCCAGGCTGGAAATGCA 3321
QY 252 ATGGCATGATCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCT 311
DB 3322 GTGGCATGATCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCT 3321
QY 312 CAATCTTCCAGTAGCTGGGACTACAGGAATGAGCTGGCCAGCTGGCTGGTTTGTG 371
DB 3382 CAGCTCCCGAGTAGTGGGACTACAGGAGTGGCTGGCCAGCTGGCTGGTTTGTG 3441
QY 372 TTTTTTAA-----TTTGGAGCCAGTGGAGTGGGCCATATCTG 412
DB 3442 TTTTTTAAAGTGACATAAAATAAATGCTTTGGGCGAGCGGAGTGGCTGGCTGGTA 3501
QY 413 ATCCAGCACTTTGGGAGCAAGGAGCGGATTAATCTTGGAGTTCAGGACTTCAAGACCA 472
DB 3502 ATCCAGCACTTTGGGAGCGCAAGTGGGTGAAT--CATGAGTTCAGGACTTCAGACCA 3559
QY 473 GCAGGCCACATGTTAAACCATGTC 499
DB 3560 ACCTGGCCAAACATGGTGAACCCCTGTC 3586

RESULT 9
US-09-764-872-518/c
; Sequence 518, Application US/09764872
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Publication No. US20030050231A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: P1125
; CURRENT APPLICATION NUMBER: US/09/764,872
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult P1125 or file wrapper
; NUMBER OF SEQ ID NOS: 957
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 518
; LENGTH: 32152
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-872-518

Query Match 33.2%; Score 165.8; DB 11; Length 32152;
Best Local Similarity 74.6%; Pred. No. 4.6e-32;
Matches 244; Conservative 0; Mismatches 62; Indels 21; Gaps 2;

QY 192 TATTTATTTATTTATTTATTTCTAGATGGAGTCTGCTGTGGCCAGGCTGGAAATGCA 251
DB 29892 TGTTTTTTTTTTTTTTTTTTTTGGAGCGGAGTCTGCTGTGGCCAGGCTGGAAATGCA 28832
QY 252 ATGGCATGATCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCT 311
DB 28931 GTGGCATGATCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCT 28772
QY 312 CAATCTTCCAGTAGCTGGGACTACAGGAATGAGCTGGCCAGCTGGCTGGTTTGTG 371
DB 29771 CAGCTCCCGAGTAGTGGGACTACAGGAGTGGCTGGCCAGCTGGCTGGTTTGTG 28712
QY 372 TTTTTTAA-----TTTGGAGCCAGTGGAGTGGGCCATATCTG 412
DB 28711 TTTTTTAAAGTGACATAAAATAAATGCTTTGGGCGAGCGGAGTGGCTGGCTGGTA 28652
QY 413 ATCCAGCACTTTGGGAGCAAGGAGCGGATTAATCTTGGAGTTCAGGACTTCAAGACCA 472
DB 28651 ATCCAGCACTTTGGGAGCGCAAGTGGGTGAAT--CATGAGTTCAGGACTTCAGACCA 28594
QY 473 GCAGGCCACATGTTAAACCATGTC 499
DB 28593 ACCTGGCCAAACATGGTGAACCCCTGTC 28547

RESULT 10
US-10-072-349-328
; Sequence 328, Application US/10072349
; Publication No. US20030054420A1;
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: P11101
; CURRENT APPLICATION NUMBER: US/10/072,349
; CURRENT FILING DATE: 2002-02-11
; Prior application removed - See file wrapper or Palm
; NUMBER OF SEQ ID NOS: 334
; SOFTWARE: PatentIn Ver. 3.1
; SEQ ID NO 328
; LENGTH: 32152
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-072-349-328

Query Match 33.2%; Score 165.8; DB 14; Length 32152;
Best Local Similarity 74.6%; Pred. No. 4.6e-32;
Matches 244; Conservative 0; Mismatches 62; Indels 21; Gaps 2;

QY 192 TATTTATTTATTTATTTATTTCTAGATGGAGTCTGCTGTGGCCAGGCTGGAAATGCA 251
DB 3262 TGTTTTTTTTTTTTTTTTTTTTGGAGCGGAGTCTGCTGTGGCCAGGCTGGAAATGCA 3321
QY 252 ATGGCATGATCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCTCAGTCT 311
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DN nucleic - nucleic search, using sw model:

Run on: October 24, 2003, 18:12:56 (Search time 1044.61 Seconds
(without alignments)
11633.309 Million cell updates/sec)

Title: US-09-830-902-1_COPY_1_500

Perfect score: 500

Sequence: 2 taactgactctgctgtgttt.....aacatggtaaacacatctgcg 500

Scoring table: IDENTITY_NJC

Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562764

Minimum DS seq length: 0

Maximum DS seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:

1: em_estha1

2: em_esthum

3: em_estint

4: em_estant

5: em_estcov

6: em_estro1

7: em_estro2

8: em_estro3

9: gb_est1

10: gb_est2

11: gb_est3

12: gb_est4

13: gb_est5

14: gb_est6

15: em_estfur

16: em_estrom

17: em_gss_hum

18: em_gss_hum

19: em_gss_pln

20: em_gss_vit

21: em_gss_fun

22: em_gss_mam

23: em_gss_mus

24: em_gss_pro

25: em_gss_rnd

26: em_gss_phg

27: em_gss_vit

28: gb_gss1

29: gb_gss2

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
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C 2	171.2	34.2	371	9	AW821209	AW821209 PM2-ST030
C 3	171.2	34.2	376	9	AW821105	AW821105 PM2-ST030
C 4	165.6	33.1	1027	10	BG622059	BG622059 6C264665

5	165.4	33.1	439	12	B1467370	B1467370 IC3le02.x
6	164.2	32.8	824	28	AC988705	AC988705 HS-3149.A
7	163.4	32.7	359	13	BQ532356	BQ532356 IJ3-H7061
8	161.8	32.4	473	3	AA433390	AA433390 zw87a09.1
9	157.9	31.6	681	29	AGC59079	AGC59079 Pan trogl
10	157.2	31.4	445	9	AA573381	AA573381 rns3a04.s
11	157.2	31.2	391	13	BQ574362	BQ574362 UI-H-E21-
12	150.8	30.2	699	12	BX990649	BX990649 UI-H-E20-
13	148.8	29.8	347	14	CE266516	CE266516 IC3E422.H
14	148.8	29.6	661	28	AQ539566	AQ539566 RPT-11-3
15	148.2	29.6	523	28	B69012	B69012 CIT-HSP-202
16	147.8	29.6	595	29	AG016569	AG016569 Homo sapi
17	146.6	29.3	416	13	B3683269	B3683269 UI-CF-CU1
18	146.6	29.3	418	13	B3675484	B3675484 UI-CF-CU1
19	146.6	29.3	104	13	BX407955	BX407955 BX47955
20	146	29.2	418	10	BF931715	BF931715 CM2-N7016
21	145.6	29.1	317	10	BE156651	BE156651 QV3-H7036
22	145	29.0	367	12	BM682825	BM682825 UI-E-E31-
23	145	29.0	436	9	AA649183	AA649183 ns3a11.s
24	144.6	28.9	492	28	B45413	B45413 HS-1361-A1-
25	144.2	28.8	627	29	AC990681	AC990681 Pan trogl
26	143.8	28.8	479	9	A1863180	A1863180 t244c04.x
27	143.6	28.7	1052	13	BX463038	BX463038 BX463038
28	143	28.6	324	12	BM982363	BM982363 UI-CF-BN1
29	142.8	28.6	353	9	A1815608	A1815608 au43b01.Y
30	142.6	28.5	441	2	HS085930	BX493132 Homo sapi
31	142.6	28.5	814	13	B0852129	B0852129 AGENCOURT
32	142.4	28.5	502	2	HS071632	BX491445 Homo sapi
33	142.2	28.4	475	9	A1361912	BX493132 Homo sapi
34	141.9	28.4	231	9	A1335357	A1361912 BK1905.x
35	141.8	28.4	639	9	AW163068	A1335357 DKFZ7628
36	141.8	28.4	729	9	AV703572	AW163068 au9jh05.Y
37	141.6	28.3	711	29	AG011827	AV703572 AV703572
38	141.2	28.2	508	12	BX991594	AG011827 Homo sapi
39	141.2	28.2	737	29	B2611164	BX991594 UI-H-D51-
40	140.6	28.1	360	9	AA847961	B2611164 WHA87567F
41	140.6	28.1	843	13	B0520971	AA847961 OD72011.S
42	140.4	28.1	404	14	C2251403	B0520971 AGENCOURT
43	140.2	28.1	190	9	A1418927	C2251403 AGENCOURT
44	139.8	28.0	827	12	B1457063	A1418927 t552101.x
45	139.8	28.0	1379	11	B0222265	B1457063 601:85984
						B0222265 Homo sapi

ALIGNMENTS

RESULT 1
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LOCUS
DEFINITION
A1343123
Repetitive element ; mRNA sequence.
A1343123.1 GI:4080329
Similar to contains Alu repetitive element; contains element PTR5

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

A1343123
tb04e07.x2 NCI-CGAP_202F Homo sapiens cDNA clone IMAGE:2052612.3
Similar to contains Alu repetitive element; contains element PTR5
Repetitive element ; mRNA sequence.
A1343123
A1343123.1 GI:4080329
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Accession: NC01343123
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs@mail.nih.gov
cDNA Library Preparation: David B. Krizman, Ph.D.
cDNA Library Arrayed by: I.M.A.G.E. Consortium, LLNL
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bic.lln.gov/abbr/image/image.html
Insert length: 437 Std Error: 0.03

Simpson, A.J.
Shotgun sequencing of the human transcriptome with OPP expressed
sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (1), 3471-3496 (2000)
200202663
MEDLINE
PUBMED
10737800
COMMENT
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL:
http://www.ludwig.org.br/scripts/gethtml2.pl?l=612=PM2-ST303-221
299-302-h054c3-1999-12-25&t4=1
Seq primer: puc 18 forward
High quality sequence start: 16
High quality sequence stop: 376.
Location/Qualifiers
1..376
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="adult"
/clone_lib="S70303"
/notes="Organ: Stomach; Vector: puc18; Site 1: Small;
Site 2: Small; A mini-library was made by cloning products
derived from CRISTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."
BASE COUNT 99 a 92 c 97 g 88 t
ORIGIN

Query Match 34.2% Score 171.27 DB 9: Length 376;
Best Local Similarity 75.3% Pred. No. 1.2e+01;
Matches 226; Conservative 0; Mismatches 73; Indels 1; Gaps 1;
CY 200 TTATTTTATTTCTAGATGAGCTTGTCTCTTCCGAGAGTGGATGGATGGATG 259
DB 363 TTATTTTATTTTACAGAGGAGCTTGTCTCTTCCGAGAGTGGATGGATGGATG 304
CY 260 ATCTCAGCTCAGTGCACCTGCACCTTTCGGTTTCAGGCAATCTCTCTCAGATGTC 319
DB 303 GTATAGTCTAGTGCACCTTGCACCTTTCGGTTTCAGGCAATCTCTCTCAGATGTC 344
CY 320 CAAGTACTGGGACTACAGAAAGAGCTGGGAGAGCTGGGAGAGCTGGGAGAGCTG 399
DB 243 CAAGTCTGGGACTACAGAAAGAGCTGGGAGAGCTGGGAGAGCTGGGAGAGCTG 485
CY 380 TTTTGGGCGAGGTGAGTGGGCGCATCTGTGATCCAGAGCTTGGGAGAGCTGGG 439
DB 184 TTTTTCCTTGGGCGAGTGGGCGCATCTGTGATCCAGAGCTTGGGAGAGCTGGG 425
CY 440 GGCGGATTTAGTGGTGGGAGTGGGAGTGGGAGTGGGAGTGGGAGTGGGAGTGGG 499
DB 124 GGCAGATCACTGAGGTGAGATGAGATGAGATGAGATGAGATGAGATGAGATGAG 65

RESULT 4
BG622059/c
LOCUS BG622059
DEFINITION BG622059.1 GI:13673430
ACCESSION BG622059
VERSION BG622059
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
1027 bp mRNA linear EST 18-APR-2001
602646665F1 NIH_MGC_79 Homo sapiens cDNA clone IMAGE:4769049 5',
mRNA sequence.
B622059

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: graphs@rail.nih.gov
Tissue Procurement: CLONTECH Laboratories, Inc.
cDNA Library Preparation: CLONTECH Laboratories, Inc.
cDNA Library Arrayed by: The I.V.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.V.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: J5CMI63; row: b column: 18
High quality sequence stop: 444.
Location/Qualifiers
1..1027
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_lib="IMAGE:4769049"
/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NIH MGC 79"
/notes="Organ: placenta; Vector: pDNP-LIB (Clontech);
Site 1: SfiI (ggccctcgcc); Site 2: SfiI (ggccctcgcc)
1: 57 and 3' adaptors were used in cloning as follows:
adaptor sequence: 5'-ATTTCAGAGCGAGCGCGGACATG-3' and 3' adaptor
(where B = A, C, or G and N = A, C, G, or T). Average
insert size 1.3 kb (range 0.5-4.0 kb); 15/15 colonies
contained inserts by PCR. This library was enriched for
full-length clones and was constructed by Clontech
Laboratories (Palo Alto, CA). Note: this is a NIH_MGC
Library."
BASE COUNT 336 a 231 c 312 g 142 t
ORIGIN

Query Match 33.1% Score 165.67 DB 10: Length 1027;
Best Local Similarity 71.4% Pred. No. 8.5e-02;
Matches 226; Conservative 0; Mismatches 79; Indels 3; Gaps 1;
CY 186 TGTGTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTT 245
DB 444 TTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTT 385
CY 246 ATSCATGTCGATGATCATCATCTCAGTGCACCTGCAGCTTTCGGTTTCAGAGATTCCTC 304
DB 394 ATTCAGTGGCAGATTTTGTTCACGTGCATTCACATCCCGGGTCAAGGATCTTC 325
CY 306 CTGCTCAATCTTCCAAAGTAGTGGAGTACAGAGATGAGTGCATCCAGCTGGT 345
DB 324 CTGCTCAGCTCCGAAATAGCTGGGATACAGAGTGCATCCAGCTGGT 268
CY 366 TGTGTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTT 425
DB 267 TGTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTT 258
CY 426 GGGAGACCAAGGAGGAGGAGGAGTACTTTCAGGTGAGGAGTTCAGACGAGGAGGAGCAT 485
DB 207 GGGAGGCTGAGGAGGAGGAGTACTTTCAGGTGAGGAGTTCAGACGAGGAGGAGCAT 448
CY 456 GGTAAAC 493
DB 147 GATGAAC 140

RESULT 5
B1467370
LOCUS B1467370
DEFINITION B1467370
ACCESSION B1467370
439 bp mRNA linear EST 22-AUG-2001
B2467370
B2467370 HR85 islet Homo sapiens cDNA 3', mRNA sequence.

The following repetitive elements were found in this cDNA sequence: 12-168, >ALU (matched complement): 139-418, >ALU 334-623, >ALU 634-661, >AT richLow_complexity
Seq primer: M13 FORWARD
POLYA=yes

FEATURES

source
1..699
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5862756"
/tissue_type="lung Focal Fibrosis"
/dev_stage="Adult"
/lab_host="CHIC3 (Life Technologies)"
/lab_libs="NCI CGAP D10"
/note="Organ: Lung; Vector: pT73-Pac (Pharmacia) with a modified polylinker; Site 1: EcoR I; Site 2: Not I; NCI CGAP D10 is a cDNA library containing the following tissue(s): A pool of lung focal fibrosis. The library was constructed according to Bonaldi, Lennon and Soares, Genome Research, 6:79-86, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT73-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is ATACGCGGTC.
TAG_D10=UI-H-D10
TAG_TISSUE=lung with fibrosis
TAG_SEQ=ATACGCGGTC

BASE COUNT 189 a 161 c 170 g 168 t 1 others
ORIGIN

Query Match 30.2%; Score 150.8; DB 12; Length 699;
Best Local Similarity 70.2%; Pred. No. 3.2e+03;
Matches 219; Conservative 0; Mismatches 87; Indels 6; Gaps 1;
QY 194 TTTTATTTTATTTTCTAGATGGAGTCTGTCTGTTGCCAGGCTGGAATGCAAT 253
DB 1 TTTTATTTTATTTTCTAGATGGAGTCTGTCTGTTGCCAGGCTGGAATGCAAT 253
QY 254 GGCATGATCTAGCTCAGTCCAACTCCACCTTCGGGTTCAAGCAATCTCTCTCTCA 313
DB 61 GGCATGATCAGATTCAGTCCAGCTCAACTCCATCTCTCAAGCAATCTCTCTCTCA 120
QY 314 ATCTTCCAGTAGCTGGAGTACAGGAATGAGCTCCGCACTGCTGCTGTTG----- 367
DB 121 GTCTCTGAGTAGTAGGACTTCAGGCAGCTGCCACATGCTGGCCACTAAGAAAAAT 180
QY 368 TTGTGTTTAAATTTTGGGCGAGTGCAGTGGGCCATATCTGTGATCCAGGCTTTGG 427
DB 181 ATTTTAAAAAATAACAGTGGTGGTACAGTGGCTCAGCGCTGTAAATCCAGCACTTGG 240
QY 428 GAGACCAAGGAGGCGGATTAATCTTCTAGCTCAGGAGTTCAAGACCGAGCCCAAGCTGG 487
DB 241 GAGGCGGAGGAGGCTGGATCACTTTAGGACACAGCTTTTGAAGCCAGCTGCGCCAACTGG 300
QY 498 TAAACCAATGTC 499
DB 301 CATACCTTGTCT 312

RESULT 13
CB266516
LOCUS
DEFINITION
1005422 Human Fat Cell 5'-Stretch Plus cDNA Library Homo sapiens
CB266516
cDNA 5', mRNA sequence.
ACCESSION
CB266516
VERSION
CB266516.1 GI:28441102
KEYWORDS
EST.
SOURCE
Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 347)
AUTHORS
Yang, R.-Z., Shuldiner, A. and Gong, D.-W.
TITLE
EST analysis of human adipose gene expression
JOURNAL
Unpublished
COMMENT
Contact: Gong Da-Wei
Division of Endocrinology, Diabetes and Nutrition
University of Maryland
660 Redwood St. #H497, Baltimore, MD 21201, USA
Tel: 410 706 1672
Fax: 410 706 1622
Email: dgong@medicine.umaryland.edu
PCR PRIMERS
FORWARD: CTCGGGAAGCGCGCCATTGTGTGGT
BACKWARD: AATAGACTCAGTATAGGGGAATGG
Seq primer: GTGGTACCGGGAATTC.

FEATURES
source
1..347
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/sex="Male and Female"
/tissue_type="Adipose"
/clone_libs="Human Fat Cell 5'-Stretch Plus cDNA Library"
/note="Vector: lambdaTriplEx"
BASE COUNT 88 a 84 c 80 g 95 t
ORIGIN

Query Match 20.8%; Score 145.8; DB 14; Length 347;
Best Local Similarity 70.7%; Pred. No. 5.9e+03;
Matches 212; Conservative 0; Mismatches 87; Indels 1; Gaps 1;
QY 194 TTTTATTTTATTTTCTAGATGGAGTCTGTCTGTTGCCAGGCTGGAATGCAAT 253
DB 24 TTTTATTTTATTTTCTAGATGGAGTCTGTCTGTTGCCAGGCTGGAATGCAAT 253
QY 254 GGCATGATCTAGCTCAGTCCAACTCCACCTTCGGGTTCAAGCAATCTCTCTCTCA 313
DB 84 GGCATGATCTAGCTCAGTCCAACTCCACCTTCGGGTTCAAGCAATCTCTCTCTCA 143
QY 314 ATCTTCCAGTAGCTGGAGTACAGGAATGAGCTCCGCACTGCTGCTGTTGTTGTTGTT 373
DB 144 GCTTCCGAGTAGTGGAGTAAAGACACTCTCTACCCCTTTTATTTAAATAGAAA 202
QY 374 TTTAAATTTTGAAGGCGAGTGGAGTGGCCATATCTGTGATCCAGCACTTTGGGAGACC 433
DB 203 CATAACTGTAAAGCCAGGCGCATGGTGGTATATGCTGTAGTCCAGCACTTTGGGAGGCT 262
QY 434 AAGGCGAGGCGGATTACTTGAGGTGAGGAGTTCAAGACCGAGCCCAACATGGTAAAC 493
DB 263 GAGTGGGGAAGATTACTTGAAGCCAGGAATTCAGACAGGCTGGGCAACATAGCAAGAC 322

RESULT 14
AQ39566/c
LOCUS
DEFINITION
RPCI-11-338N10.TV RPCI-11 Homo sapiens genomic clone RPCI-11-338N10
genomic survey sequence.
ACCESSION
AQ39566
VERSION
AQ39566.1 GI:4870140
KEYWORDS
GSS.
SOURCE
Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 661)
AUTHORS
Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter
J.C.
TITLE
Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building
JOURNAL
Unpublished

Query Match 20.8%; Score 145.8; DB 14; Length 347;
Best Local Similarity 70.7%; Pred. No. 5.9e+03;
Matches 212; Conservative 0; Mismatches 87; Indels 1; Gaps 1;
QY 194 TTTTATTTTATTTTCTAGATGGAGTCTGTCTGTTGCCAGGCTGGAATGCAAT 253
DB 24 TTTTATTTTATTTTCTAGATGGAGTCTGTCTGTTGCCAGGCTGGAATGCAAT 253
QY 254 GGCATGATCTAGCTCAGTCCAACTCCACCTTCGGGTTCAAGCAATCTCTCTCTCA 313
DB 84 GGCATGATCTAGCTCAGTCCAACTCCACCTTCGGGTTCAAGCAATCTCTCTCTCA 143
QY 314 ATCTTCCAGTAGCTGGAGTACAGGAATGAGCTCCGCACTGCTGCTGTTGTTGTTGTT 373
DB 144 GCTTCCGAGTAGTGGAGTAAAGACACTCTCTACCCCTTTTATTTAAATAGAAA 202
QY 374 TTTAAATTTTGAAGGCGAGTGGAGTGGCCATATCTGTGATCCAGCACTTTGGGAGACC 433
DB 203 CATAACTGTAAAGCCAGGCGCATGGTGGTATATGCTGTAGTCCAGCACTTTGGGAGGCT 262
QY 434 AAGGCGAGGCGGATTACTTGAGGTGAGGAGTTCAAGACCGAGCCCAACATGGTAAAC 493
DB 263 GAGTGGGGAAGATTACTTGAAGCCAGGAATTCAGACAGGCTGGGCAACATAGCAAGAC 322

RESULT 14
AQ39566/c
LOCUS
DEFINITION
RPCI-11-338N10.TV RPCI-11 Homo sapiens genomic clone RPCI-11-338N10
genomic survey sequence.
ACCESSION
AQ39566
VERSION
AQ39566.1 GI:4870140
KEYWORDS
GSS.
SOURCE
Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 661)
AUTHORS
Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter
J.C.
TITLE
Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building
JOURNAL
Unpublished

COMMENT

Other GSSs: RPCI-11-338N10.TJ
 Contact: Shaying Zhao, William Niernan, Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbs@tigr.org
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pictor de Jong
 (pictor@igmc.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
 Research Genet cs (info@resgen.com). BAC end search page:
 http://www.tigr.org/tcdb/hungen/bac_end_search.html.
 Seq primer: 17
 Class: BAC ends.

FEATURES

source
 1..661
 /organism="Homo sapiens"
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 /db_xref="GB:7629729"
 /db_xref="taxon:9606"
 /clone="RPCI-11-338N10"
 /sex="Male"
 /cell_type="JY-phocytes"
 /clone_lib="RPCI-11"
 /note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
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 BASE COUNT 141 a 202 c 148 g 170 t
 ORIGIN

Query Match 29.8%; Score 149.8; DS 26; Length 661;
 Best Local Similarity 68.9%; Pred. No. 3.6e+03;
 Matches 219; Conservative 0; Mismatches 97; Indels 2; Gaps 1;
 QY 184 TGTGTTTATTTATTTATTTATTTATTTCTAGATGGAGTGTGCTGTGTTGGCAGGCT 243
 |||||
 DB 358 TGTGTTTATTTCTATTTATTTATTTCTAGATGGAGTGTGCTGTGTTGGCAGGCT 299
 QY 244 GGAATGCATGGCATGATCTAGCTCAGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTC 263
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 DB 298 GAGGCCAGTGGTGGCATCGGCTGATCAGGCCAGGCTCGAACGCCGCTGAGGCGATCC 239
 QY 304 TGTGCTCAATGTTCCAAATGAGCTGGGACTACAGATGAGCTGCGGCACTGTCCT-- 261
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 DB 238 TGTGCTCGATGCTCCAGATGCTGGGATACAGCGTGGATGAGCGATGATGAGCTAA 233
 QY 362 GGTGTTGTTGTTTTMAATTTTGAAGCCAGGTGGCATGTCGCGCATATCTGTGATCCAGCA 421
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 DB 178 ATTATGTTTAAAGAGGCTCAGTCGGGCTGGCTGTGCTTACACCTGTAAATCCAGAA 119
 QY 422 GTTGGGAGCAGGAGCAGGCGGATCTACTTGGAGTCGAGGTCGAGGTCGAGGAGCAGGCGCA 481
 |||||
 DB 118 GTTGGGAGGCGAAGCGAGGAGATCACTTGAGATCAGGAGTTCAGAGCAGGCTGACCA 59
 QY 482 ACATGGTAAACCATGTC 499
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 DB 58 ACATGGTAAACCATGTC 41

RESULT 15

B69012 521 bp DNA linear GSS 21-JUN-1995
 LOCUS CIT-HSP-2025G21.TF CIT-HSP Homo sapiens genomic clone 2025G21,
 DEFINITION genomic survey sequence.

ACCESSION B69012

VERSION B69012.1 GI:2667722

KEYWORDS GSS.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 523)

AUTHORS

Adams, V.D., Rounsley, S.D., Field, C.E., Bass, S., Linher, K., Golden,
 X., Berry, K., Granger, D., Suh, E., Wible, C., Shizuya, H., Simon, X.,
 and Venter, J.C.
 Use of a random BAC End Sequence Database for Sequence-Ready Map
 Building

TITLE

Unpublished

JOURNAL

Other GSSs: CIT-HSP-2025G21.TF

COMMENT

Contact: Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: mdamas@tigr.org
 Clones are available from Research Genetics (info@resgen.com). BAC
 end search page:
 http://www.tigr.org/tcdb/hungen/bac_end_search.html.
 Seq primer: M13-21
 Class: BAC ends.

FEATURES

source
 1..523
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 /mol_type="genomic DNA"
 /db_xref="GB:7646951"
 /db_xref="taxon:9606"
 /clone="2025G21"
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 /cell_type="Sperm"
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 /note="Vector: pBeloBAC11; Site_1: HindIII; Site_2:
 HindIII"

BASE COUNT 127 a 127 c 134 g 135 t
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 Best Local Similarity 72.7%; Pred. No. 4.6e+03;
 Matches 232; Conservative 0; Mismatches 83; Indels 4; Gaps 3;

QY 182 TTGTTGTTTATTTATTTATTTATTTCTAGATGGAGTGTGCTGTGTTGGCAGG 241
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 DB 31 TTTTCTCTGTGAATTTTGTGTTGTTTGAAGCAAGGCTGTGCTGTGTTGCCCAAG 90
 QY 242 CTGGAATGCATGGCATGATCTAGCTCAGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTC 300
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 DB 91 CGGAGTCAGTAGCATGATCATAGCTCAGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTC 150
 QY 301 TCTCTCTGCTCATCTTCCAGTAGCTGGGATCAGGATGAGCTCCGGCAGCTGGCC 360
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 DB 151 TCTCTCTGCTCAGCTTCTCAAGTAGCTAGGACCAAGCATGGGCCATCACACCT-AAC 209
 QY 361 TGGTTTGTGTTTAAATTTTGAAGCCAGGTGAGTGGCCCATATCTGTGATCCAGC 420
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 DB 210 TATTTTATTTTATTTTATTTAGTAGAGCCAGGCGGAGTGGCTTACGCTGTATTCGAGC 269
 QY 421 ACTTTGGAGACCAAGCGAGGCGGATTTACTTGGTTCAGGATTCAGACCGACGAGGCG 480
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 DB 220 ACTTTGGAGCGCAAGCGCGGTGGAT--CATGAGTCAGGAGTTCAGACCGAGCTGGTC 327
 QY 481 ACATGGTAAACCATGTC 499
 |||||
 DB 328 ACATGGTAAACCATGTC 346

Search completed: October 24, 2003, 22:50:57

Job time : 1049.8 secs

GenCore version 3.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

CM nucleic - nucleic search, using sw model

Run on: October 24, 2003, 15:43:05 ; Search time 134.09 seconds
(without alignments)
14808.112 Million cell updates/sec

Title: US-09-830-902-1_COPY_4000_4500

Perfect score: 501
Sequence: 1 ccaagctgttgattacag.....tgaatgaagctagctagcc 501

Scoring table: IDENTITY NUC
Gapop 100, Gapext 1.0

Searched: 2888711 seqs, 2045441386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

GenEmbl:

1: gb_ba:

2: gb_hg:

3: gb_in:

4: gb_or:

5: gb_ov:

6: gb_pat:

7: gb_ph:

8: gb_pl:

9: gb_pr:

10: gb_ro:

11: gb_sts:

12: gb_sy:

13: gb_un:

14: gb_vit:

15: em_ba:

16: em_fun:

17: em_hum:

18: em_in:

19: em_mu:

20: em_or:

21: em_ov:

22: em_pat:

23: em_ph:

24: em_pi:

25: em_pl:

26: em_ro:

27: em_sts:

28: em_un:

29: em_vit:

30: em_hg_hum:

31: em_hg_inv:

32: em_hg_other:

33: em_hg_mus:

34: em_hg_pln:

35: em_hg_rod:

36: em_hg_sam:

37: em_hg_vit:

38: em_sy:

39: em_hg_hum:

40: em_hg_mus:

41: em_hg_other:

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	501	100.0	101584	9	CNS01B55	AL121655 BAC seque
2	501	100.0	110000	6	AX093471	AX093471 Sequence
3	501	100.0	110000	6	WSA24603	AL246003 Homo sapi
4	501	100.0	155943	9	AC012364	AC012364 Homo sapi
5	98.8	19.7	127192	9	HS036502	AL035462 Human DNA
6	97.4	19.4	73036	2	AC091063	AC091063 Homo sapi
7	94	18.8	194835	9	AL158158	AL158158 Human DNA
8	93.6	18.7	142728	9	HSJ792G4	AL159163 Human DNA
9	85	17.0	75946	9	EX248112	EX248112 Human DNA
10	85	17.0	153770	9	AL353911	AL353911 Human DNA
11	85	17.0	187035	2	EX388586	EX388586 Homo sapi
12	84.4	16.8	63409	2	AC022992	AC022992 Homo sapi
13	84.4	16.8	97811	9	AC008634	AC008634 Homo sapi
14	84.4	16.6	104475	9	AC011394	AC011394 Homo sapi
15	84.2	16.6	121394	2	AL137799	AL137799 Homo sapi
16	84.2	16.8	190570	2	AL157894	AL157894 Homo sapi
17	83.8	16.7	93418	2	AC068379	AC068379 Homo sapi
18	83.8	16.7	179887	2	AC069489	AC069489 Homo sapi
19	83.8	16.7	198952	2	AP002016	AP002016 Homo sapi
20	83.4	16.6	61840	2	AC130291	AC130291 Homo sapi
21	83.4	16.6	132383	9	EX296545	EX296545 Human DNA
22	83.4	16.6	197676	2	AC022544	AC022544 Homo sapi
23	82.5	16.5	126613	9	HSJ603114	AL122001 Human DNA
24	82.2	16.4	163709	2	AP001155	AP001155 Homo sapi
25	82.2	16.4	185228	9	AC023043	AC023043 Homo sapi
26	82.2	16.4	195489	2	AP001159	AP001159 Homo sapi
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ALIGNMENTS

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ACCESSION AL121655
VERSION AL121655.1 GI:5002398
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SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 101584)
AUTHORS Hazan, J., Focknechten, N., Mavel, D., Paternotte, C., Sanson, D.,
Ariague, F., Davoine, C.S., Cruaud, C., Durif, A., Wincker, P.,

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutera; Primates; Caccarhini; Hominiidae; Homo.

REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES

Weissenbach, J. and Hazar, J.
Cloning, expression and characterization of the sp4 gene responsible for the most frequent form of autosomal spastic paraplegia
Patent: WO 01/8193-A 1 15-MAR-2001
CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (CNRS) (FR)
Location/Qualifiers
1. 110000
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ACCESSION AJ246003
VERSION AJ246003.1 G1:6273492
KEYWORDS Spast gene; spastin protein; SPG4-linked hereditary spastic paraplegia.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Hazan, J., Fokknechten, N., Mavel, D., Paternotte, C., Samson, D., Arizguenave, P., Davoine, C.S., Gruaud, C., Durr, A., Wincker, P., Brattier, P., Catolico, L., Barbe, V., Burgunder, J.M., Prud'homme, C.F., Brice, A., Fontaine, B., Prud'homme, C., Weissbach, C., Durr, A. and Hazan, J.
TITLE Spastin, a new AAA protein, is altered in the most frequent form of autosomal dominant spastic paraplegia
JOURNAL Nat. Genet. 23 (3), 296-303 (1999);
MEDLINE 20255425
PUBMED 10610178
REFERENCE 2
AUTHORS Fokknechten, N., Mavel, D., Byrne, P., Davoine, C., Gruaud, C., Boetsch, D., Samson, D., Coutinho, P., Hurchinson, V., McMonagle, P., Burgunder, J., Tartaglione, A., Heinzel, C., Fokknechten, P., Paternotte, C., Brice, A., Fontaine, B., Prud'homme, C., Weissbach, C., Durr, A. and Hazan, J.
TITLE Spectrum of SPG4 mutations in autosomal dominant spastic paraplegia
JOURNAL Hum. Mol. Genet. 9 (4), 637-644 (2000)
MEDLINE 20164302
PUBMED 10699187
REFERENCE 3
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-JUN-1999) Genoscope, Genoscope - Centre National de Sequencage, BP 191, Evry 91006, FRANCE
COMMENT E-mail : seqref@genoscope.cns.fr - Web : www.genoscope.cns.fr. The sequence is the result of the assembly of 2 BAC clones: R-335p14 and 563N4, respectively from RPCI-11 and C12A_976_SKB library.
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gene

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NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., third quality >= 30). An attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RP2311 human BAC library was made from the blood of one male donor as described by Georgerakopoulos, Wong, P. Y., Haug, J., Paterno, M., Caraceni, J. and de Jong, P. (1993). An improved approach to construction of bacterial artificial chromosomes. *Biotechniques* 15: 63-67. The clone may be obtained either from Genomics, Inc. (<http://www.genomics.com>) or from the Roswell Park Cancer Institute and coworkers at the Roswell Park Cancer Institute (<http://bacc.med.buffalo.edu>).

VECTOR: 3BACE3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP1-43P79; the clone sequenced to the right is RP1-78E13. Actual start of this clone is at base position 1 of RP1-44D15; actual end is at base position 5543 of RP1-44D15.

Data from AC01232 and AC00991 was used to finish this clone, AC01234. The sequence fidelity between bases 11265 to 11263 can not be guaranteed due to an unresolved homopolymeric run. The sequence between 11245 to 11263 is single stranded.

538

[illegible]

REFERENCE
AUTHORS

2 (bases 1 to 73096)
 Birrer, B., Litton, L., Nusbaum, C., Lander, E., Allen, M., Anderson, S.,
 Barna, N., Bastien, V., Boguslavsky, L., Boekhauser, B., Brown, A.,
 Camarata, J., Campopiano, A., Chang, J., Choepel, Y., Colangelo, M.,
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 Zaitoun, C., Zerbek, S., Zimmer, A. and Zody, M.

TITLE
JOURNAL

Direct Submission
 Submitted (25-MAR-2003) Whitehead Institute/MIT Center for Genome
 Research, 120 Charles Street, Cambridge, MA 02141, USA

COMMENT

All repeats were identified using RepeatMasker:
 Smith, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WBR

Web site: <http://www.seq.wi.mit.edu>

Contact: sequence_submission@genome.wi.mit.edu

----- Project Information

Center Project Name: 112746

Center Clone Name: 1078_M_23

NOTE: This record contains 88 individual
 sequencing reads that have not been assembled into
 contigs. Runs of N are used to separate the reads
 and the order in which they appear is completely
 arbitrary. Low-pass sequence sampling is useful for
 identifying clones that may be gene-rich and allows
 overlap relationships among clones to be deduced.
 However, it should not be assumed that this clone
 will be sequenced to completion. In the event that
 the record is updated, the accession number will
 be preserved.

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DB 33923 AGGTGACACCTTTGCAATATAACTGTGATCTTCTGAATAAGAGATGTAGTCTATT 33864  
  
QY 270 TCCATATATTGCACTAGAGTTGGCTTCTGACTTGTCTTGCAGATGGAATGTAGTACAA 329  
DB 33963 CCCATCCCTTGAACCTGGATAGGCTTTGTGACTTTTTCGACCATGTAGGTACAGCAA 33804  
  
QY 330 ATGACACTGTGCAACTTTGGATTTTAGGTTTCGAGAGAACTTACACTTCCACTCACACT 389  
DB 33803 GTATATTGTTAGGACTTACAAATCTAGACCCCAAGAAAGCTTGTAGCTTCCATTCTCACC 33744  
  
QY 390 CTCTTGAACACAGATGCAATGTAAAGAGTCAGGCTATCTCTCTAGAG 439  
DB 33743 CTCTTGAACAC---TGCCCTGAAGAAGCTCAGTCTCACTACTGAG 33657  
  
RESULT 8  
HSDJ792G4/c
```

LOCUS HSDJ792G4 142722 bp DNA linear PRI 31-DEC-2003
DEFINITION Human DNA sequence from clone RP4-792G4 on chromosome RP1.2-32.2. Contains a transcription factor gene, a pseudogene similar to tetraatricopeptide protein, ESTs, CA repeat (C15438), STSs, GSSs and CpG islands, complete sequence.
ACCESSION AL049636
VERSION G1:11367799
KEYWORDS HTG; CUS438.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 142722):
AUTHORS Bkaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE Howden P.
JOURNAL Direct Submission
COMMENT Submitted (20-DEC-2003): Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: turquerry@sanger.ac.uk
 requests: clonerequests@sanger.ac.uk
 On Dec 22, 2003 this sequence version replaced GI:5531522.
 During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.
 This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated repeat sequence elements. Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em: EMBL; SW: SWISSPROT; Tr: TrEMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/EGP/Chr1>
 RP4-792G4 is from the library RPi-4 constructed by the group of Pletier de Jong. For further details see <http://www.chori.org/bacpac/home.htm>
FEATURES VECTER: pCYPAC2
 This sequence is the entire insert of clone RP4-792G4.
 Location/Qualifiers
 1..142726
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosomes="1"
 /map="p1.2-32.2"
 /clone="RP4-792G4"
 /clone_lib="RPi-4"
 repeat_region 1..142722 repeat: matches 1..62 of consensus"
 repeat_region 1724..1797
 /note="Char1e4 repeat: matches 1233..1355 of consensus"
 repeat_region 1799..1842
 /note="22 copies 2 mer at 84% conserved"
 repeat_region 2387..2414
 /note="14 copies 2 mer to 89% conserved"
 misc_feature 2036..10523
 /note="match: GSS: EM:AO165153"
 repeat_region 13213..13497
 /note="AUS01 repeat: matches 1..309 of consensus"
 repeat_region 13619..13930
 /note="AUS01 repeat: matches 1..308 of consensus"
 repeat_region 14743..17426
 /note="L1M1 repeat: matches 879..1556 of consensus"
 repeat_region 17806..17891
 /note="L1M1 repeat: matches 1547..1625 of consensus"
 repeat_region 13885..19497
 /note="L1M1 repeat: matches 1623..1761 of consensus"

23778..24183
 /note="match: GSS: EM:AO270799"
 repeat_region 24773..24820
 /note="24 copies 2 mer at 81% conserved"
 repeat_region 25736..25769
 /note="17 copies 2 mer to 82% conserved"
 repeat_region 27577..27630
 /note="27 copies 2 mer to 74% conserved"
 repeat_region 27634..27880
 /note="VER2 repeat: matches 14..255 of consensus"
 repeat_region 28476..28606
 /note="AUS0/x repeat: matches 3..133 of consensus"
 repeat_region 31013..31234
 /note="L1M1 repeat: matches 5946..6172 of consensus"
 repeat_region 31235..31323
 /note="MER44C repeat: matches 1..89 of consensus"
 repeat_region 31322..31503
 /note="MER44C repeat: matches 235..47 of consensus"
 repeat_region 31504..31670
 /note="FRAX repeat: matches 1..168 of consensus"
 repeat_region 31674..31962
 /note="MER44C repeat: matches 413..697 of consensus"
 repeat_region 31977..32132
 /note="L1M1 repeat: matches 6159..6323 of consensus"
 repeat_region 32644..32930
 /note="AUS35 repeat: matches 1..301 of consensus"
 repeat_region 33861..33937
 /note="AUSP/q repeat: matches 175..311 of consensus"
 repeat_region 34422..34632
 /note="MLT1-INTERNAL repeat: matches 828..881 of consensus"
 repeat_region 36772..36859
 /note="MLT1-INTERNAL repeat: matches 1083..1157 of consensus"
 repeat_region 40721..40819
 /note="MER33 repeat: matches 1..113 of consensus"
 repeat_region 41156..41263
 /note="MER33 repeat: matches 133..238 of consensus"
 repeat_region 41326..41485
 /note="L1M2 repeat: matches 6151..6304 of consensus"
 repeat_region 41795..41983
 /note="L1M2 repeat: matches 6041..6351 of consensus"
 repeat_region 41904..41957
 /note="XPR33 repeat: matches 243..297 of consensus"
 repeat_region 42270..42307
 /note="19 copies 2 mer aa 81% conserved"
 misc_feature 44966..44552
 /note="match: GSS: EM:AO422993"
 misc_feature 46318..46409
 /note="CpG island"
 /evidence="not experimental"
 complement(5650)..47101
 /note="match: GSS: EM:AZ15581"
 52353..53789
 /gene="d079204.1"
 52353..53789
 /gene="d079204.1"
 /note="match: CDNAS: EM:AF052249 Em:U39843 Em:J13220 Em:J59832 Em:AC71554 Em:U04198 Em:X0460 Em:AF052251 Em:U11690 Em:AF104932 Em:U37272 Em:J87393 Em:X92592 Em:X95603 Em:L38607 Em:U13202 Em:L13201 Em:X93291 Em:L22760 Em:AB001572 Em:X93128 Em:S79041 Em:AJ011652 Em:AF023915 Em:X74143 Em:U90538 Em:X55955 Em:S82462 Em:AF172889 Em:L12142 Em:L12143 Em:M14511 Em:L12703 Em:J13222 Em:AB028021
 match: ESTs: EM:AI739311 Em:AI066718 Em:AI554075 Em:AA248965 Em:AI391745 Em:AI323408
 match: proteins: Sw:P79772 Sw:Q61560 Sw:Q16676 Sw:P11087 Sw:Q61345 Sw:Q64732 Sw:P55318 Sw:P22452 Tr:Q93613 Sw:P22453 Sw:Q01149 Tr:Q51415 Sw:P26791 Sw:P26312 Tr:Q22751 Tr:Q3784 Sw:Q99958 Sw:Q98917 Sw:Q12950 Sw:Q63244 Tr:Q70220 Sw:P06924 Sw:Q63245 Sw:Q64733 Tr:Q86778 Tr:Q73779 Wp:CB07013 Sw:P55787 Sw:Q60688


```

18187 AATTGAGATGCTATACGGTCGATTTGTTCCAGAGACGACTGCACACTGCATCTA 18246
204 TCTCATATATATGCCCTTTGCAATGATGATTTTCTACTGCTCTATATACAGATGCGAG 263
18247 TC-----TTCATGCGCTTTGCAATATGACTTTTCCACTTTCCATCCAGACATAGACT 18301
264 TATTTTCCATATATTCGACTAGAGTTGGCTCTCTACTGCTTTGTA-CAATGGAAATGT 328
19302 CTATCTTACTCCCTTAAATCTGGACTAATCTATGAGTTGCTTTAACCAGATATCATGT 18361
323 ACTGAGAAATGACACTGTGCAACTTTGGATTTCAGTTTCAGAGAGACTTACAGTTCCAC 352
18362 GGTAGAAATGACAAATGTGCAACTTCAGAGAGAACTTCAAGATTCGCCACCTTCAT 18421
393 TCACACTCTCTTGGAAACGAGATGCAATG 411
18422 TTTTATGCTTGGAAATGCAAGTGCATG 18450

RESULT 10
AL953911 Human DNA sequence from clone RP11-242D:2 on chromosome 9, complete
sequence.
LOCUS AL953911
DEFINITION AL953911.4 GI:25788317
VERSION HTG.
KEYWORDS Homo sapiens (human);
SOURCE Homo sapiens
ORGANISM Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 150770)
Direct Submission
Submitted (13-DEC-2002) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquerry@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Dec 13, 2002 this sequence version replaced gi:25788317.
***** Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquerry@sanger.ac.uk
*****
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e. phred quality >=
30). An attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest, except on the rare
occasion of the clone being a YAC.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases.
Ent, EMBL; Swi, SWISSPROT; Tr, TrEMBL; Wp, WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/Celegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr9
RP11-242D12 is from the library RPC1-11.1 constructed by the group
of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACe3.6.
Location/Qualifiers

```

```

source
1. 150770
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="9"
/clone="RP11-242D12"
/clone_1b="RPC1-11.1"
BASE COUNT 45356 a 30392 c 30483 g 44540 t
ORIGIN
Query Match 17.34; Score 95; DB 9; Length 150770;
Best Local Similarity 62.5%; Pred. No. 7e-11;
Matches 158; Conservative 0; Mismatches 95; Indels 6; Gaps 2;
QY 144 ATTTCACATATATCTGTCTAAATTTACTTGCAGAGATGGCCACAAATTCCTCTTA 203
DB 96018 AATTGAGATGCTCTATACGGTCGATTTTCCAGAGACGACTGCACACTGCATCTA 96077
QY 204 TCCATCATATATGCCCTTTGCAATGCTGACTTTGCTTCTCTTCAAGATGTGGAGC 263
DB 96078 TC-----TTCATGCGCTTTTTCAGATATGACTTTTCCACTTTCCATCCAGACATAGACT 96132
QY 264 TATTTTCCATATATTCGACTAGAGTTGGCTTCTGACTTGGCTTCTGACTTGGTTTGA-CAATGGAAATGT 352
DB 96133 CTATCTTACTCCCTTAAATCTGGACTAATCTATGAGTTGCTTTTACCAATATCATGT 96192
QY 323 AGTACAATGACACTGTGCAACTTTGGATTTCAGTTTCAGAGAGACTTACAGTTCCAC 392
DB 96193 GGTAGAAATGACAAATGTGCAACTTCAGAGAGAACTTCAAGATTCGCCACCTTCAT 96252
QY 393 TCACACTCTCTTGGAAACGAGATGCAATG 411
DB 96253 TTTTATGCTTGGAAATGCAAGTGCATG 9628;

RESULT 11
BX088588 Homo sapiens chromosome 9 clone RP11-5019, *** SEQUENCING IN
LOCUS PROGRESS ***
DEFINITION BX088588
VERSION BX088588.4 GI:2780504K
KEYWORDS HTG; HTGS; PHASE2
SOURCE Homo sapiens (human);
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 187035)
Direct Submission
Submitted (07-APR-2003) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquerry@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Jan 20, 2003 this sequence version replaced gi:27803285.
***** Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquerry@sanger.ac.uk
***** Project Information
Center project name: BA5019
***** Summary Statistics
Assembly program: XGAP4; version 4.5
Chemistry: Dye-terminator; 10% cf reads
Consensus quality: 18703 bases at least Q40
Consensus quality: 187031 bases at least Q80
Consensus quality: 187034 bases at least Q20
Insert size: 187035; sum-of-contigs
Quality score: 196616; 2.8% error; agarose-fp
Quality coverage: 16.5% in Q20 bases; sum-of-contigs Quality
coverage: 15.8% in Q20 bases; agarose-fp
*****
* NOTE: This is a 'working draft' sequence. It currently

```

- * consists of 1 contigs. Gaps between the contigs
- * are represented as runs of N. The order of the pieces
- * is believed to be correct as given, however the sizes
- * of the gaps between them are based on estimates that have
- * provided by the submitter.
- * This sequence will be replaced
- * by the finished sequence as soon as it is available and
- * the accession number will be preserved.
- * i 187035: contig of 187035 bp in length.

Location/Qualifiers

i 187035
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="9"
 /clone="RP11-5013"
 /clone_lib="RPC-11.1"

misc_feature
 i 187035
 /notes="assembly fragment 0:450"
 BASE COUNT 55353 a 37202 c 37514 g 56966 t
 ORIGIN

Query Match 17.0% Score 95; CB 2; Length 187035;
 Best Local Similarity 62.5%; Pred. No. 6.8e-11;
 Matches 166; Conservative 3; Mismatches 95; Indels 6; Gaps 2;
 QY 144 ATTCTACATATATGCTGCTAAATTACTTSCAAAGATGCGCACAGCAATTCCTCTA 203
 DB 125354 AATTGAAGATGCTCATACGGTCGATGCTTCCAAAGACGACTGCAACACGCTTCTA 125295
 QY 204 TCGTCATATATGCGCTTGGAAATGTCACCTTGTCTTCTTCTATCATGAGATGCGAGC 263
 DB 125294 TC-----TTACATGCGCTTTGGCAATGATGCTTTTCATTTTCATCAAGACATAGACT 125240
 QY 264 TCAATTTCCCATATATGCACTAGATGTCGCTTGTGCTGCTTTGA CATGGAATGT 322
 DB 125239 CTAATTTACTTCCCTTAATATCGGACTATATGAGTGTCTTACCAATATCATGT 125180
 QY 323 AGTACAATGACACTGTGCACTTTTGATTTTGGTTTTCGAGAGACTTACCTTCAC 382
 DB 125179 GGTAGAATGACATGTGCAACTTCCAAAGACAAAGACTTCAAGATCCCACTTCAT 125120
 QY 383 TCACACTCTCTTGGAAACCAAGATGCAATG 41;
 DB 125119 TTTTATGCTTGGAAATGCACTGCGCATG 125091

RESULT 12
 AC022982/c
 LOCUS
 DEFINITION Homo sapiens chromosome 18 clone RP11-545P23 map 18, LOW-PASS
 SEQUENCE SAMPLING.
 AC022982
 AC022982.2 GI:9163727
 VERSION
 KEYWORDS HTG; HTGS PHASE0.
 SOURCE Homo sapiens (human)
 ORGANISM
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE
 1 (bases 1 to 63409)
 Birren, B., Linton, L., Nussbaum, C., Lander, E., Abraham, H., Allen, N.,
 Anderson, S., Baldwin, J., Barna, N., Becker, R., Bedalov, E.,
 Boguslavsky, L., Bookshgaiter, B., Brown, A., Buckner, G., Castle, A.,
 Choquet, Y., Colangelo, X., Collins, S., Collymore, A., Cooke, P.,
 DeRubeis, K., Dewar, K., Domero, M., Doyle, M., Fanestor, J.,
 Ferrell, P., FitzHugh, W., Forrest, C., Gage, D., Galagan, J.,
 Gardyna, S., Grant, G., Haggis, B., Heaford, A., Horton, L.,
 Howland, J. C., Johnson, A., Jones, C., Karn, L., Kavatas, A., Klein, C.,
 Landers, T., Lechoczky, J., Levine, R., Liu, C., Liu, J., Locke, K.,

Macdonald, P., Marquis, N., McEwan, P., McGuck, A., McKerran, K.,
 McPherson, R., Meadim, J., Meneus, J., Morrow, J., Naylor, J.,
 Norman, C. H., O'Connor, T., O'Donnell, P., Oliver, T. M., Peterson, K.,
 Pierre, N., Pisani, C., Pollara, V., Raymond, C., Riley, R., Rothman, D.,
 Roy, A., Santos, R., Severy, P., Spencer, B., Stange, Thomas, N.,
 Stojanovic, N., Subramanian, A., Talamas, A., Tesfaye, S., Theodore, J.,
 Tittel, A., Vassiliev, H., Viel, R., Vo, A., Wu, X., Wyman, D., Ye, W.,
 Zimmer, A. and Zody, J.

TITLE
JOURNAL

COMMENT

Submitted 107-FEB-2003; Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Jul 13, 2003 this sequence version replaced GI:9163727.
 All repeats were identified using RepeatMasker.

Smith, A.P.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

Center: Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIER
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submission@genome.wi.mit.edu
 Project information
 Project name: L694
 Center project name: 545_P_23

- * NOTE: This record contains 73 individual
- * sequencing reads that have not been assembled into
- * contigs. Runs of N are used to separate the reads
- * and the order in which they appear is completely
- * arbitrary. Low-pass sequence sampling is useful for
- * identifying clones that may be gene-rich and allows
- * overlap relationships among clones to be deduced.
- * However, it should not be assumed that this clone
- * will be sequenced to completion. In the event that
- * the record is updated, the accession number will
- * be preserved.

758: contig of 758 bp in length
 759 858: gap of 100 bp
 859 1633: contig of 766 bp in length
 1634 1733: gap of 100 bp
 1734 2499: contig of 766 bp in length
 2500 2599: gap of 100 bp
 2600 3355: contig of 756 bp in length
 3356 3455: gap of 100 bp
 3456 4210: contig of 755 bp in length
 4211 4310: gap of 100 bp
 4311 5031: contig of 721 bp in length
 5032 5131: gap of 100 bp
 5132 5902: contig of 771 bp in length
 5903 6023: gap of 100 bp
 6024 6781: contig of 779 bp in length
 6782 6881: gap of 100 bp
 6882 7661: contig of 780 bp in length
 7662 7761: gap of 100 bp
 7762 8519: contig of 758 bp in length
 8520 8619: gap of 100 bp
 8620 9396: contig of 777 bp in length
 9397 9496: gap of 100 bp
 9497 10268: contig of 772 bp in length
 10269 10368: gap of 100 bp
 10369 11144: contig of 776 bp in length
 11145 12144: gap of 100 bp
 12145 12115: gap of 100 bp
 12116 12582: contig of 767 bp in length
 12583 12982: gap of 100 bp
 12983 13754: contig of 772 bp in length
 13755 14624: contig of 773 bp in length
 14625 14724: gap of 100 bp
 14725 15482: contig of 758 bp in length
 15483 15582: gap of 100 bp
 15583 16316: contig of 754 bp in length
 16317 16416: gap of 100 bp

JOURNAL Submitted (07-DEC-2001) DOE Joint Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA

COMMENT On Dec 7, 2001, this sequence version replaced gi:11752565.
Draft Sequence Produced by DOE Joint Genome Institute
www.jgi.doe.gov
Finishing Completed at Stanford Human Genome Center
www.shgc.stanford.edu
Quality: Phrap Quality >=40 100% of Sequence:
Estimated Total Number of Errors is 0.
Location/Qualifiers
1. 104475
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTB-143D11"
PAGE COUNT 31967 a 10178 c 19721 g 13609 t
ORIGIN

Query Match 16.6%; Score 84.4; DB 9; Length 104475;
Best Local Similarity 62.6%; Pred. No. 1e-10; 2; Gaps 2;
Matches 164; Conservative 0; Mismatches 96; Indels 2;

QY 138 AAGGAAATCTACACATATCAATCTTGCTAAATTAATCTTGCAAAGATGCCACACAAATTC 137
DB 94402 AAGTCTATTATTACTCAGATGAGAGTGTGTTTACCAAGGTGSCCAGACAAATTA 94343

QY 198 CTCCTATCTCATATATATGCCCCCTTSCAATGTGACCTTTCCTACTTCTCTATCAAGATG 257
DB 94342 TCCCATCCCTATATCCGAGGACCTTGCTGTGTGACTCCACCCTCTCTCAAGAG-GA 94284

QY 258 TGGAGCTATTCTCCCATATATTGCACATAGATTGGCTTCTGACTTCCTTGA-CAATG 316
DB 94283 AAGAGTCAATTTATTTCCTCTGSAATTAAGCTGCTGTGATTCGTTTACCAATA 94284

QY 317 GAATGTAGTACAAATGACACTGTGCAACTTGGATTTTAGTTTCGAGAGAACTTACACC 376
DB 94221 GAAGGTGTACAGATGACACAGCAACTTGTGGCCCTAGGTCTCAAGAGCATCTCAAC 94164

QY 377 TCCCACTCACCTCTCTGGAA 395
DB 94163 ATCTGTTTACGCCCAATGGAA 94142

RESULT 15
LOCUS AL137799
DEFINITION Homo sapiens chromosome 1 clone RP5-1199A8, *** SEQUENCING IN
PROGRESS ***, 6 unordered pieces.
ACCESSION AL137799
VERSION AL137799.4 G:10039491
KEYWORDS RTG; RTGS PHASE1; RTGS_CANCELLED.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1
Plumb, B.
Direct Submission
Submitted (09-JUL-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk Clone
requests: clonerequests@sanger.ac.uk
On Sep 8, 2000 this sequence version replaced gi:1211917.
----- Genome Center
Center: Sanger Centre
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
----- Project Information
Center project name: d01199A8
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Sequencing vector: plasmid; L08752; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads
 Consensus quality: 119318 bases at least Q40
 Consensus quality: 119965 bases at least Q30
 Consensus quality: 120403 bases at least Q20
 Insert size: 120394; sum-of-contigs
 Quality coverage: 128726; 9.8% error; acetone-fp
 Quality coverage: 6.06x in Q20 bases; sum-of-contigs Quality
 coverage: 5.79x in Q20 bases; acetone-fp

- NOTE: This is a 'working draft' sequence. It currently consists of 6 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

- 54341: contig of 54341 bp in length
- 54441: gap of 100 bp
- 60268: contig of 5827 bp in length
- 60369: gap of 100 bp
- 90782: contig of 30414 bp in length
- 90783: gap of 100 bp
- 90883: contig of 11860 bp in length
- 102742: gap of 100 bp
- 102743: contig of 100 bp
- 102943: contig of 4442 bp in length
- 107285: gap of 100 bp
- 107385: contig of 14010 bp in length.

FEATURES

Location/Qualifiers

1..121394
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="1"
 /clone="RP25-1193A8"
 /clone_lib="RPCL-5"

1..54341
 /note="assembly fragment:00965"
 fragment_chain:1
 clone_end:576
 vector_side:left
 54442..60268
 /note="assembly fragment:01389"
 fragment_chain:1
 60369..90782
 /note="assembly fragment:01706"
 fragment_chain:1
 90883..102742
 /note="assembly fragment:00399"
 fragment_chain:2
 102843..107284
 /note="assembly fragment:00472"
 fragment_chain:2
 107385..121394
 /note="assembly fragment:01054"
 fragment_chain:2

BASE COUNT 31229 a 31246 C 25603 g 28816 t 500 others

ORIGIN

Query Match 16.8%; Score 84.2; DB 2; Length 121394;
 Best Local Similarity 65.7%; Freq. No. 1.1e-10;
 Matches 153; Conservative 0; Mismatches 78; Indels 2; Gaps 2;

QY 216 TGGCCCTTGGCAATGTGACATTTGGTACTTCTCT-ATCAGAGATGGAGGCTTATTTCCCA 274
 |||||
 DB 119265 TGTCCAGTAGACATGTGACTTTGCTGTTCTGTCATTAGAGATGAGTCTGTTTCCCA 119324
 |||||
 QY 275 TATATGCACTAGAGTTGGGCTTCTGACTTGGCTTGA-CAATGGAATGATGACAAATGA 133
 |||||
 DB 119325 CCTCTGAATCTGGCTGATCTATGATTTGTTTGGACCAGTAAAGTCTGGTGAAGGGA 119384
 |||||
 QY 334 CACTGTGCAACTTTGGATTTAGGTTTCGAGAGAACTTACCGTTCCACTCAGCTCTCT 393
 |||||

DB 119385 TACTGTGCAACTTCTGCATGTAGGGCTCAAGAGGCTTCGCAACTTCCACGGTCACTTTGG 119444
 QY 394 TGGAAACCCAGATGGCAATGTAAAGAGTCAAGGCTATCTCTGTAGAGACATATG 446
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 DB 119445 AGCTCTTCTGGGGCTCTGTGAGAGAGGCTTGGACCAACCTGCTTGGAGACATGTG 119497
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Search completed: October 24, 2003, 21:06:02
 Job time : 1389.29 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.
OM nucleic acid nucleic search, using sw model
Run on: October 24, 2003, 15:39:20 / Search time 12f 25f 25f Seconds
/ without alignment
/ 1072.159 Million Cell updates/sec

Title: US-09-830-902-1_COPY_4000_4500
Perfect score: 501
Sequence: 1 ccaagtcgctggatcag.....tgaatgagcctagctagcctaggcc 501

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 2552756 seqs, 134979217 residues
Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: N_Geneseq_10.unclust.*
1: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA1880.DAT.*
2: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA1881.DAT.*
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4: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA1883.DAT.*
5: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA1884.DAT.*
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21: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA1900.DAT.*
22: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*
25: /SIDSI/gcgdata/geneseq/geneseq-emb1/NA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	501	100.0	110000	22	AAF84800 Nucleotide sequence
C 2	82.8	16.5	502	20	AAZ22641 Novel CDNA sequence
C 3	82.6	16.5	854	20	AAZ22642 Novel nucleotide s
C 4	82.6	16.5	27859	22	ABA19635 Human immune syst
C 5	82.6	16.5	27869	22	AAK66517 Human immune/haema
C 6	70	14.0	218336	25	ABQ76678 Androgen receptor
C 7	64.6	12.9	5239	24	ABU32273 Human immune syste
C 8	63	12.6	1241	21	AAC74337 Human secreted pro

3	51.4	12.3	1473	22	AAC05322 Human secreted pro
10	41.4	12.3	1454	22	AAC05372 Human G protein-co
11	61	22.2	67	22	AAZ45921 CDNA encoding nove
12	61	22.2	670	24	ABK41689 Human immune syste
13	60.4	12.1	5239	24	ABU32272 EST clone G51. H
C 14	60.2	12.0	651	20	AAV88447 Human CDNA sequenc
C 15	59.4	11.5	2477	22	AAH75555 N. meningitidis pa
16	56.8	11.3	1040	21	AAZ81696 Human chromosome 3
17	56.6	11.3	567571	25	ABU53224 CDNA encoding huma
18	56.4	12.3	1777	25	ABX12157 Human immune/haema
C 19	56.4	12.3	5434	22	AAK76220 Human immune/haema
20	53.8	10.7	3535	22	AAK65228 Human immune/haema
21	53.8	10.7	3535	22	AAK73493 Human polynucleoti
22	53.6	10.7	12.4	22	AAI59283 Human polynucleoti
23	53.6	10.7	1275	22	AAI61569 Human protein kina
24	53.6	10.7	1586	21	AAZ66793 Human protein kina
25	53.6	10.7	1586	22	ADL11344 Human protein kina
26	53.6	10.7	1586	24	ABQ76287 Human protein kina
27	53.6	10.7	1868	24	AAI17048 Human protein kina
28	53.6	10.7	1868	25	ABX14972 Human protein kina
C 29	53.6	10.7	2142	22	AAH42264 Cell cycle protein
C 30	53.6	10.7	2375	22	AAI01478 Human kinase PKN-
31	53.6	10.7	2539	22	AAH76219 Human TGF-beta rec
32	53.6	10.7	3240	22	AAI75345 Novel protein kina
33	53.6	10.7	3387	22	AAI44687 Human CDNA cffere
34	53.2	10.6	90220	24	ABX83576 Human transglutari
C 35	53.2	10.6	100000	24	ABQ74541 Human transglutari
C 36	52.6	10.5	37658	25	ABQ8210 Cell cycle protein
37	52.2	10.4	2141	22	AAH42263 Human secreted pro
38	51.6	10.3	245	21	AAI31581 Human digestive sy
39	51.6	10.3	1434	22	AAK89569 Human colorectal c
40	51.6	10.3	1434	22	AAI57826 Genomic DNA # enc
41	51.6	10.3	1434	24	ABX99803 Human protease exp
42	51	10.2	502	23	ABV5821 Human immune syst
C 43	50.4	10.1	10137	22	ABX18984 Human immune/haema
C 44	50.2	10.0	389	22	AAK64465 Human immune/haema
C 45	50.2	10.0	972	22	AAK73575 Human immune/haema

ALIGNMENTS

RESULT 1	AAF84800	AAF84800 standard; DNA: 110000 bp.
XX	AAI949500:	
AC	AAI949500:	
XX	09-JUL-2002 (first entry)	
XX		Nucleotide sequence of the human SPQ4 gene.
DE		Human: SPQ4 gene; spastin; PSF-AD; gene therapy;
KW		autosomal dominant familial spastic paraplegia; ss.
XX		Homo sapiens.
XX		Location/Qualifiers
FT	Key	9932..10203
FT	CDs	/tag= a
FT	exon	/note= "contains introns"
FT		9932..10471
FT		/tag= b
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Db 4300 ACTTGCTTTGACAAATGSAATGTAGTACAAATGACAGCTGTGCACTTTAGATT 4309
 CC
 QY 361 CAGAGACATTACACTTCCACTGCACTCTCTGGGAAACAGAGATGATGAAGAAGT 429
 CC
 Db 4360 CAGAGACATTACACTTCCACTGCACTCTCTGGGAAACAGAGATGATGAAGAAGT 4410
 CC
 QY 421 CAGGGCTATCTGCTAGACAGATATGCTCCAGCTATATGAGCATGACATCTGACAT 480
 CC
 Db 4420 CAGGGCTATCTGCTAGACAGATATGCTCCAGCTATATGAGCATGACATCTGACAT 4479
 CC
 QY 481 ATGAATGAGGCTAGTAGGCC 501
 CC
 Db 4480 ATGAATGAGGCTAGTAGGCC 4500
 CC

RESULT 2

AAZ22642/c

ID AAZ22642 standard; cDNA; 502 BP.

XX

AC AAZ22641;

XX

DT 08-DEC-1999 (first entry)

XX

DE Novel cDNA sequence 1 derived from the b2HFLS20W cDNA library.

XX

KW interferon; novel sequence; pathogenic infection; antiviral; cell fate;

XX

KW cell proliferation; immune disorder; leukemia; wound healing; antibody;

XX

KW assay; ds.

XX

CS Homo sapiens.

XX

PM WO9945117-A2.

XX

PM WO9945117-A2.

XX

PM 10-SEP-1999.

XX

PM 04-MAR-1999; 99WO-US03727.

XX

PM 24-MAR-1999; 98US-0034878.

XX

PM 23-AUG-1998; 98US-0137348.

XX

PM (HYSE-1) HYSEQ INC.

XX

PM Drmanac RT, Crkvenjakov R, Dickson M, Drmanac S, Labat I;

XX

PM Leshkowitz D, Kita D, Ford C, Mulero CJ;

XX

PM WPI; 1999-551041/46.

XX

PM New interferon polynucleotides useful for treating disorders involving

XX

PM pathogenic infection, cell fate and differentiation, inflammation etc.

XX

PM Claim 1; Fig 2; 94pp; English.

XX

PM This a novel sequence isolated from the b2HFLS20W cDNA library. prepared

XX

PM from human fetal liver-stem tissue, with another novel sequence

XX

PM (AAZ22642). The polypeptide sequences corresponding to these nucleic

XX

PM acid sequences have a striking homology to the interferon family of

XX

PM polypeptides.

XX

PM The interferon polypeptides may be administered therapeutically to

XX

PM treat/prevent disorders involving pathogenic infection (e.g. viral

XX

PM and bacterial infections); or cell fate and differentiation (e.g. immune

XX

PM disorders, leukemias, wound healing, tissue growth etc.). Interferons

XX

PM are known to have antiviral activity and are believed to be involved in

XX

PM regulatory processes such as cell proliferation and differentiation.

CC The polynucleotides are also useful to detect polynucleotides encoding
 CC the polypeptides e.g. to diagnose disorders for susceptibility to
 CC disorders relating to polypeptide expression, or mark polypeptide-
 CC expressing tissues. They may also be used to produce probes useful to
 CC detect related sequences or for gene mapping.
 XX

XX Sequence 502 BP; 141 A; 97 C; 143 G; 118 T; 3 other;

XX Query Match 16.5%; Score 92.8; DB 20; Length 502;

XX Best Local Similarity 63.8%; Pred. No. 2.1e-14;

XX Matches 157; Conservative 0; Mismatches 87; Indels 2; Gaps 2;

QY 203 ATCTCATATATAGCCCTTGCATATGACATTTGCTTCTCTATCTCT-ATCAAGATGGA 261

Db 347 ATCTCCCATCTGCTCCAGTACATGACATTTCTCTCTCTCAITTAAGAGATGA 288

QY 282 GCTTATTTCCATATATGACATAGATTTGCGCTTCTGACTTGTCTTGA-CMAATGGAT 320

Db 287 GTCGTTTTCCCACTCTTGAATCTGGCTGATCTTATGATTTGTTTGACCAATAGT 328

QY 321 GTAGTACAAATGACATCTGCAACTTTGGATTTTAGTCTTTGAGAGAACTTACACCTTCC 380

Db 227 CTGGTGGAGGGATGCTGCGCACTTCTGCATGTAGGCTCAAGAAGCCTCGCACTTCC 368

QY 331 ACTGACACTCTCTTGGAAACGAGATGCAATGTAAGAAATCAGGCTATCTGCTAGAGA 440

Db 167 ACCTGCACTTTTGGAGCTCTTCTGCGCTCTGTGAAGAAGCCTGGCAACCTCTGAGA 408

QY 441 CATATG 446

Db 107 CATGTG 102

RESULT 3

AAZ22642/c

ID AAZ22642 standard; cDNA; 854 BP.

XX

AC AAZ22642;

XX

DT 08-DEC-1999 (first entry)

XX

DE Novel nucleotide sequence IFN-HY from the b2HFLS20W cDNA library.

XX

KW interferon; novel sequence; pathogenic infection; antiviral; cell fate;

XX

KW cell proliferation; immune disorder; leukemia; wound healing; antibody;

XX

KW assay; ds.

XX

CS Homo sapiens.

XX

PM WO9945117-A2.

XX

PM 10-SEP-1999.

XX

PM 04-MAR-1999; 99WO-US03727.

XX

PM 04-MAR-1999; 98US-0034878.

XX

PM 23-AUG-1998; 98US-0137348.

XX

PM (HYSE-1) HYSEQ INC.

XX

PM Drmanac RT, Crkvenjakov R, Dickson M, Drmanac S, Labat I;

XX

PM Leshkowitz D, Kita D, Ford C, Mulero CJ;

XX

PM WPI; 1999-551041/46.

XX

PM New interferon polynucleotides useful for treating disorders involving

XX

PM pathogenic infection, cell fate and differentiation, inflammation etc.

XX

PM Claim 1; Fig 2; 94pp; English.

XX

PM This a novel sequence isolated from the b2HFLS20W cDNA library. prepared

CC from human fetal liver-spleen tissue, with another novel sequence
CC (AA222641). The polypeptide sequences corresponding to these nucleic
CC acid sequences have a striking homology to the interferon family of
CC polypeptides and this nucleotide sequence has been designated IFN-HV.
CC The interferon polypeptides may be administered therapeutically to
CC treat/prevent disorders involving pathogenic infection (e.g. viral
CC and bacterial infections) or cell fate and differentiation (e.g. immune
CC disorders, leukemias, wound healing, tissue growth etc.). Interferons
CC are known to have antiviral activity and are believed to be involved in
CC regulatory processes such as cell proliferation and differentiation.
CC The polypeptides may also inhibit platelet aggregation and therefore be
CC used to treat/prevent conditions involving inflammatory responses. They
CC can be combined with a carrier in compositions which can be administered
CC to treat disorders as above, and used to generate antibodies useful in
CC polypeptide detection and purification. They can also be used to
CC identify binding compounds, by contacting with a compound and detecting
CC complex formation, optionally within a cell in which the complex drives
CC expression of a reporter gene. They (and the polynucleotides) are
CC therefore useful in drug screening assays e.g. for the above disorders.
CC The polynucleotides are also useful to detect polynucleotides encoding
CC the polypeptides e.g. to diagnose disorders of susceptibility to
CC disorders relating to polypeptide expression, or mark polypeptide-
CC expressing tissues. They may also be used to produce probes useful to
CC detect related sequences or for gene mapping.
XX
SQ Sequence 854 BP: 251 A; 180 C; 243 G; 190 T; 3 other.

Query Match 16.51; Score 82.6; DB 20; Length 854;
Best Local Similarity 65.24; Pred. No. 3e-14;
Matches 152; Conservative 0; Mismatches 79; Indels 2; Gaps 2;

QY 216 TGCCCTTGGCAATGAGCTTGGTACTTCTCT-ATCAGATGTGGAGCTATTTCGCA 274
DB 334 TGCCAGTACATGATGATTTGCTCTCTCTCTATAGAGATGAGCTCTTCCGCA 275

QY 275 TATATTCAGTACAGTGGCTTCTGACTTGTCTTGA-CAATGGAAATGATACAAATGA 331
DB 274 CCTCTTGAATCGGGCTGATCTATGATTTGTTGACCGGTAAGTCTGGTGGAGGGA 215

QY 334 CACTGTGCACTTGGATTTTAGGTTTCAGAGAACTTACACTTCCACTCAGCTCTCT 393
DB 214 TGGTGGCACTTCTGATGATGAGGCTCAGGCTCAGAGAGCTCCAGCTCATTTCG 155

QY 394 TGAACACGATGCAATGATAGAGAGCTCAGGCTATCTCTGCTAGAGAGATATG 446
DB 154 ASCCTTTCGGGCTCTGTGAGAGAGCTTGGAGAGAGCTTGGAGAGATATG 102

RESULT 4
ABA19635
ID ABA19635 standard; DNA; 27863 BP.
XX
AC ABA19635;
XX
DT 23-JAN-2002 (first entry)
XX
DE Human nervous system related polynucleotide SEQ ID NO 11966.
XX
KW Human; neutropic; neuroprotective; cytostatic; dermatological; virucide;
KW immunosuppressive; antiinflammatory; anti-HIV; antibacterial; vulnery;
KW antiparkinsonian; antiskilling; antianemic; antiarthritic; cancer;
KW antirheumatic; hepatotropic; cerebrotropic; antineoplastic; antiinflammatory;
KW antiallergic; antidiabetic; antitumor; anticonvulsant; antifungal;
KW antiparasitic; cardiac; immune disorder; cardiovascular disorder;
KW neurological disease; infection; nephrotropic; gene therapy; vaccine; ds.
XX
OS Homo sapiens.
XX
PN NC0200159061-A2.
XX
PD 16-AUG-2001.
XX
PF 17-JAN-2001; 2001MO-US01334.

XX 31-JAN-2003; 2000US-0179365.
PR 24-FEB-2003; 2000US-0180628.
PR 24-FEB-2003; 2000US-0184564.
PR 22-MAR-2003; 2000US-0186350.
PR 16-MAR-2003; 2000US-0189874.
PR 17-MAR-2003; 2000US-0190376.
PR 18-APR-2003; 2000US-0198123.
PR 19-MAY-2003; 2000US-0205215.
PR 27-JUN-2003; 2000US-0209467.
PR 28-JUN-2003; 2000US-0214886.
PR 30-JUN-2003; 2000US-0215135.
PR 27-JUL-2003; 2000US-0216547.
PR 27-JUL-2003; 2000US-0216880.
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PR 14-AUG-2003; 2000US-0224118.
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PR 14-AUG-2003; 2000US-0225757.
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PR 18-AUG-2003; 2000US-0226279.
PR 22-AUG-2003; 2000US-0226681.
PR 22-AUG-2003; 2000US-0226868.
PR 22-AUG-2003; 2000US-0227182.
PR 23-AUG-2003; 2000US-0227809.
PR 30-AUG-2003; 2000US-0228924.
PR 01-SEP-2003; 2000US-0229287.
PR 01-SEP-2003; 2000US-0229343.
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PR 05-SEP-2003; 2000US-0229509.
PR 05-SEP-2003; 2000US-0229513.
PR 06-SEP-2003; 2000US-0230437.
PR 06-SEP-2003; 2000US-0230438.
PR 08-SEP-2003; 2000US-0231242.
PR 08-SEP-2003; 2000US-0231243.
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PR 08-SEP-2003; 2000US-0231413.
PR 08-SEP-2003; 2000US-0231414.
PR 08-SEP-2003; 2000US-0232080.
PR 08-SEP-2003; 2000US-0232081.
PR 12-SEP-2003; 2000US-0231965.
PR 14-SEP-2003; 2000US-0232397.
PR 14-SEP-2003; 2000US-0232398.
PR 14-SEP-2003; 2000US-0232399.
PR 14-SEP-2003; 2000US-0232400.
PR 14-SEP-2003; 2000US-0232401.
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PR 14-SEP-2003; 2000US-0233064.
PR 14-SEP-2003; 2000US-0233065.
PR 21-SEP-2003; 2000US-0234223.
PR 21-SEP-2003; 2000US-0234274.
PR 25-SEP-2003; 2000US-0234997.
PR 25-SEP-2003; 2000US-0234998.
PR 26-SEP-2003; 2000US-0235484.
PR 27-SEP-2003; 2000US-0235834.
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PR 29-SEP-2003; 2000US-0236367.
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PR 29-SEP-2003; 2000US-0236369.
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PR 02-OCT-2000; 2000US-0246882.
PR 02-OCT-2000; 2000US-0237837.
PR 03-OCT-2000; 2000US-0237039.
PR 03-OCT-2000; 2000US-0237019.
PR 03-OCT-2000; 2000US-0237040.
PR 13-OCT-2000; 2000US-0239335.
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PR 20-OCT-2000; 2000US-0240460.
PR 20-OCT-2000; 2000US-0241763.
PR 20-OCT-2000; 2000US-0241766.
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PR 20-OCT-2000; 2000US-0241809.
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PR 20-OCT-2000; 2000US-0242221.
PR 01-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
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PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
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PR 08-NOV-2000; 2000US-0246611.
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PR 17-NOV-2000; 2000US-0249207.
PR 17-NOV-2000; 2000US-0249208.
PR 17-NOV-2000; 2000US-0249209.
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PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249243.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249264.
PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249299.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250391.
PR 01-DEC-2000; 2000US-0251160.
PR 03-DEC-2000; 2000US-0251013.
PR 03-DEC-2000; 2000US-0251988.
PR 03-DEC-2000; 2000US-0256719.
PR 06-DEC-2000; 2000US-0251479.
PR 06-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251899.
PR 08-DEC-2000; 2000US-0251992.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.
PR (HUMA-1) HUMAN GENOME SCI INC.
XX
PR Roser CA, Barash SC, Ruben SM;
XX WPI; 2001-541565/60.
XX
XX Nucleic acids encoding 3224 human nervous system antigen polypeptides,
XX useful for preventing, diagnosing and/or treating nervous system.
PT

PT cancers and metastases -
XX Disclosure; SEQ ID NO 1:966; 1701pp + Sequence Listing; English.
XX
XX The invention relates to novel genes (AB11004-AB12134) and proteins (AB14478-AB18001) useful for preventing, treating or ameliorating medical conditions e.g. by protein or gene therapy. The genes are isolated from a range of human tissues disclosed in the specification. The nucleic acids, proteins, antibodies and antagonists are useful in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast and ovarian cancer and other cancers of the adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver, lung or urogenital; (b) immune disorders e.g. Addison's disease, allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid arthritis and ulcerative colitis; (c) cardiovascular disorders such as myocardial ischaemia; (d) wound healing; (e) neurological diseases e.g. cerebral ataxia and epilepsy; and (f) infectious diseases such as viral, bacterial, fungal and parasitic infections.
XX Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pat_sequences.
XX
SQ Sequence 27869 BP; 7054 A; 6673 C; 6322 G; 7820 T; 0 other;
Query Match 16.5%; Score 82.6; DB 22; Length 27869;
Best Local Similarity 65.2%; Pred. No. 1:1e-13;
Matches 152; Conservative 0; Mismatches 79; Indels 2; Gaps 2;
QY 216 TGCCCTTTGCAATGTGACTTGTCTACTTCTCT-ATCAAGATGTGGAGCTATTTCCTCA 274
DB 27536 TGTCCAGTACAATGTGACTTTGCTGCTTCTGCTATTAGAGATGGACTTGTTCCTCCA 27525
QY 275 TATATTGCACTAGAGTGGCTTCTGACTTCTTGA-CAATGGATGTACAAATGA 332
DB 27596 CTTCTTGAATCTGGCTGATCTTATGATTTGTTGACCACTAAGCTGGGAGGGA 27655
QY 334 CACTGTGCAACTTGGATTTAGGTTTCGAGAGAACTTACACCTTCCACTCACACTCTCT 393
DB 27656 TGTGTGCAATTTCTGATGTAGGCTCAAGAGCTCGCAACTTCCAGCTCATTTTGG 27715
QY 394 TGAATCCAGATGCAATGTAAAGAGAGTCAAGGCTATCTGTAGAGACATATG 446
DB 27716 AGCTCTTCTGCGCTCTGTGAAGAGGCTGCACCAATCTGTGAGACATGTG 27768
RESULT 5
AC AAK6517/C
ID AAK6517 standard; DNA: 27869 BP.
XX AAK6517;
XX
XX Q6-NOV-2001 (first entry)
XX
XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:21329.
XX
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer; cytostatic; gene therapy; vaccine; metastasis; ds.
XX Homo sapiens.
XX WC200157182 A2.
XX 09-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-0501354.
XX
XX 31-JAN-2000; 2000US-0179065.
XX 04-FEB-2000; 2000US-0180628.
XX 24-FEB-2000; 2000US-0184664.
XX 02-MAR-2000; 2000US-0186350.
XX 16-MAR-2000; 2000US-0189874.
XX 17-MAR-2000; 2000US-0190076.
XX


```
PA (EPIC-1) EPIDEMIOLOGICS AG.
XX
XX Olek A. Piepenbrock C, Berlin K;
XX
XX WPI: 2002-130909/17.
XX
XX Nucleic acid comprising fragment of chemically modified gene, useful
XX for diagnosis and treatment of diseases associated with abnormal
XX cytosine methylation.
XX
XX Claim 1: SEQ ID NO 246; 32pp - Sequence Listing: German.
XX
XX The present invention provides a number of human immune system associated
XX genes which are modified by the methylation of cytosines. The sequences
XX can be used in the diagnosis and treatment of immune system disorders,
XX including eye diseases such as retinopathy, neovascular glaucoma and
XX macular degeneration, arteriosclerosis, anemia, cancer, acute myeloid
XX leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
XX rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel
XX diseases. The present sequence is a gene of the invention.
XX
XX Sequence 5239 BP; 1541 A; 54 C; 1235 G; 2375 T; 0 other;
SQ
Query Match 12.9%; Score 54.6; DB 24; Length 5239;
Best Local Similarity 51.3%; Pred. No. 1.2e-05;
Matches 174; Conservative 0; Mismatches 154; Indels 1; Gaps 1;
QY 62 TTTTATATTCAGCAAAATGTTGGAAATCTCTCTTAAATGAGATGCTTAAGTGGCG 121
DB 3273 TTTAATACCTTTAAAGAACACCAACCAATTTAATTAATTAATTAATTAATTAAT 3214
QY 122 TCTGACATGAGGTAGAGGAAATCTACACATATATATGCTTAATTAATTAATTAAT 13;
DB 3213 TCATTACCTTTTAAACCAACCAACCAACCAATTTAATTAATTAATTAATTAATTA 3154
QY 182 ATGGCCAGCAAAATCTCTCTCTATCATATATATGCTTTGCAATGAGATTGCTA 241
DB 3153 ATATACCATCATGATCTCTCTCTATCATATATACCTAGTACCTTACCTTACCA 3104
QY 242 CTCTCTATCAGATGCGAGCTTATTTTCCATATATGACATAGAGTTGGCTTCTGA 301
DB 3093 TTTCTCTATCAAAATTAATTTATCTCTCCATCTTAAATTTAAATCAAGCTTATA 3024
QY 302 CTCTCTTGA-CAGTGGAGATAGTACAAATGACACTGTGCACTTGGATTTGAGTIT 350
DB 3033 TTTACTTTACCAATAAATAATACAAATAATATATATATATATATATATATAT 2934
QY 361 CGAGAGAACTTACACTTCCACTCACACTCTCTCTGGAAA 399
DB 2973 CGAAAGCTTACAACTTATATTTTAAATCTCGAAAA 2935
RESULT 8
AAC74337/c
ID AAC74337 standard; cDNA: 1241 BP.
XX
XX AAC74337;
XX
XX 02-FEB-2001 (first entry)
XX
XX Human secreted protein gene 1 SEQ ID NO:11.
XX
XX Human: secreted protein; diagnosis; cytostatic; immunosuppressive;
XX neotropic; neuroprotective; antiviral; antiallergic; hepatotropic;
XX antidiabetic; antiinflammatory; antitumor; vulnervary; anticonvulsant;
XX antibacterial; antifungal; aniparasitic; cardiac; gene therapy;
XX food additive; preservative; chemosore identification; cancer;
XX female reproductive system disorder; immune disorder; wound healing;
XX cardiovascular disorder; neurological disease; infectious disease;
XX infection; ss.
XX
XX Homo sapiens.
XX
XX 17-JUL-2001 (first entry)
```

```
PK WC20058340-A2.
XX
XX CS-OCT-2000.
XX
XX 23-MAR-2000; 2000MO-US07724.
XX
XX 26-MAR-1999; 99US-C12651C.
XX 07-JAN-2000; 2000US-0174850.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Ruben SM, Komatsoulis G;
XX
XX WPI: 2000-594638/56.
XX P-PSDB; AAB39402.
XX
XX Fifty nucleic acid molecules encoding human secreted proteins, useful
XX in the prevention, treatment and diagnosis of cancer, immune disorders,
XX cardiovascular disorders and neurological diseases
XX
XX Claim 1: Page 329; 391pp; English.
XX
XX The polynucleotide sequences given in AAC74337 to AAC74336 encode the
XX human secreted proteins given in AAB39402 to AAB39451. AAB39452 to
XX AAB39484 represent human secreted polypeptide sequences and proteins
XX homologous to them, which are given in the exemplification of the present
XX invention. Human secreted proteins have activities based on the tissues
XX and cells the genes are expressed in. Example of activities include:
XX cytostatic; immunosuppressive; neotropic; neuroprotective; antiviral;
XX antiallergic; hepatotropic; antidiabetic; antiinflammatory; antitumor;
XX vulnervary; anticonvulsant; antibacterial; antifungal; antiparasitic; and
XX cardiac. The polynucleotides and polypeptides are useful for preventing,
XX treating or ameliorating a medical condition in e.g. humans, mice,
XX rabbits, goats, horses, cats, dogs, chickens or sheep. The polypeptides
XX can also be used as a food additive or preservative to increase or
XX decrease storage capabilities. The polynucleotide are useful for
XX chromosome identification. They are also useful as probes for diagnosing
XX a disorder related to the female reproductive system, particularly breast
XX and/or ovarian cancer. They are also useful in the gene therapy of breast
XX and ovarian cancer. Secreted protein nucleic acids, proteins,
XX antibodies, agonists and antagonists are useful in the diagnosis,
XX treatment and prevention of: (a) cancer; (b) immune disorders; (c)
XX cardiovascular disorders; (d) wound healing; (e) neurological diseases;
XX and (f) infectious diseases such as viral, bacterial, fungal and
XX parasitic infections. AAC74328 to AAC74336 and AAB39402 represent
XX sequences used in the exemplification of the present invention.
XX
XX Sequence 1241 BP; 363 A; 266 C; 257 G; 355 T; 0 other;
SQ
Query Match 12.6%; Score 63; DB 21; Length 1241;
Best Local Similarity 67.4%; Pred. No. 2e-08;
Matches 118; Conservative 0; Mismatches 55; Indels 2; Gaps 2;
QY 146 TCTACACATAATCATTTGCTTAATTAATTAATTAATTAATTAATTAATTAATTAAT 204
DB 803 TATTACATCTTTAATGTGGAAGTTATTAGCAAGATGCTACAATTAATTTCTTCAT 744
QY 205 CCTCAATATATATGCTTTGCAATGAGCTTGTACTT-CTCTATCAAGATGTGAGC 263
DB 743 TTCCATACATATACCATTTTATTAATATGCTCTGCCACTTGGCCCATCAAGATGGAAT 694
QY 264 TTATTTCCCATATATGCGACTAGAGTGGCTTCTGACTTGTCTTGTGACATGGA 318
DB 683 TTAATTTCCCATACCAATCAATCAGGGGTGTGTATGTAACATGCTTTGGCCCAACGA 629
RESULT 9
AAC05322
ID AAC05322 standard; cDNA: 1479 BP.
XX
XX AAC05322;
XX
XX 17-JUL-2001 (first entry)
```



```
PF 01-NOV-2000; 2000WO-US00045.
XX
XX
XX 05-NOV-1999; 99US-016359.
PR 30-JUN-2000; 2000US-0215133.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PA
XX
XX Ruben SM, Kcmatsoulis GA, Moore PA, Birse CE, Ni C;
PI
XX WPI; 2001-308778/32.
DR P-PSDB; AAE01506.
XX
XX New nucleic acid molecules encoding 28 human secreted proteins for
PT diagnosing, preventing, treating or ameliorating medical conditions and
PT used as food additives or preservatives.
XX
PS Claim 1; Page 477-478; 562pp; English.
XX
XX AAE05300-AAE05379 represent cDNAs corresponding to 28 human secreted
CC protein genes, and AAE01416-AAE01513 represent the proteins they encode.
CC AAE01514-AAE01544 represent human secreted protein fragments or variants.
CC The genes and their secreted proteins are useful for preventing,
CC treating or ameliorating medical conditions, e.g., by protein or gene
CC therapy. Pathological conditions can be diagnosed by determining the
CC amount of the new protein in a sample or by determining the presence of
CC mutations in the new genes. Specific uses are described for each of the
CC 28 genes, based on the tissues in which they are most highly expressed,
CC and include developing products for the diagnosis or treatment of
CC proliferative disorders, cancer, tumours, foetal and developmental
CC abnormalities, haematopoietic disorders, diseases of the immune system,
CC AIDS, autoimmune diseases (e.g., rheumatoid arthritis), inflammation,
CC allergies, neurological disorders (e.g., Alzheimer's disease,
CC Parkinson's disease), cognitive disorders (e.g., schizophrenia, asthma,
CC skin disorders (e.g., psoriasis), sepsis, diabetes, atherosclerosis,
CC cardiovascular disorders, oncogenic disorders, kidney disorders,
CC gastrointestinal disorders, pregnancy-related disorders, endocrine
CC disorders, and infections. The proteins can also be used to aid wound
CC healing and epithelial cell proliferation, to prevent skin aging due to
CC sunburn, to maintain organs before transplantation, for supporting cell
CC culture of primary tissues, to regenerate tissues, to identify their
CC cognate ligands or binding partners, and in chemotaxis, and can be used
CC as a food additive or preservative to modify storage properties.
CC Antibodies specific for a protein of the invention can be used in
CC alleviating symptoms associated with the disorders mentioned above, and
CC in diagnostic immunoassays e.g., radioimmunoassay or enzyme-linked
CC immunosorbent assay (ELISA). The present sequence represents a human
CC secreted protein-encoding cDNA of the invention.
XX
XX Sequence 1484 BP; 418 A; 337 C; 300 G; 422 T; 7 other;
SQ
Query Match: 12.3%; Score 61.4; DP 22; Length 1484;
Best Local Similarity 57.2%; Pred. No. 6.4e-08;
Matches 131; Conservative 0; Mismatches 96; Indels 2; Gaps 1;
QY 228 TGTGACTTTCCTCTATCAAGATGTGGAGCTTATTTTCCCATATATTCGACTAGA 369
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
632 TATGTGTTGGCCACCTCCCATCAAGACATGGTGCCTATTTTTCATCCTTTGTCTCG 691
QY 289 GTTGGCTTTCGACTTCCTTTGACATGGAATGTAGTACAAATGACACTGGCACTTTG 348
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
692 CTGGCTTGTGTTCCTATTCACCAATAGGTATGGTATGGAATGGCATTCGCACTTCT 751
QY 349 GATTTTAGGTTTCGAGAGAACTTACACTTCCACTCCACTCTCTTTGGAAACAGATGCA 408
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
752 GAGCCCAACCTT--AAAAGACATTTGAATTTTGTCTTAGTGAGAGGGGAATCAAGATACC 809
QY 409 ATGTAAGAGAGTGGGCTATCTCTCTAGACAGATATGTCCAGCTAAT 457
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
810 CTGTAAGAGAGTCTGTGCTCTTCTTGAAGACAGAGGCTCTCGAAGAT 858
RESULT 11
AA546911
```

```
1C AAS46911 standard: cDNA; 670 BP.
XX
XX AAS46911:
AC
XX
XX 19-DEC-2001 (first entry):
DT
XX
XX Human G protein-coupled receptor (GPCR) cDNA #93.
DE
XX
XX Human; G protein-coupled receptor; GPCR; mental disorder; schizophrenia;
XX neurological disorder; metabolic disorder; cancer; rheumatoid arthritis;
XX thyroid disorder; neurodegenerative disorder; cardiovascular disorder;
XX renal failure; autoimmune disorder; hyperproliferative disorder; HIV; ss;
XX human immunodeficiency virus; viral infection; neuroprotective;
XX human immunostimulant; neuroleptic; nootropic; tranquiliser; antidepressant;
XX anorectic; gene therapy.
XX
XX Homo sapiens.
OS
XX
XX NC0200160858-A2.
XX
XX 20-SEP-2001.
XX
XX 16-MAR-2001; 2001WO-US08456.
XX
XX 16-MAR-2000; 2000US-187733P.
XX 16-MAR-2000; 2000US-189977P.
XX 16-MAR-2000; 2000US-189917P.
XX 16-MAR-2000; 2000US-193916P.
XX 16-MAR-2000; 2000US-189960P.
XX 29-MAR-2000; 2000US-192155P.
XX 29-MAR-2000; 2000US-192342P.
XX 29-MAR-2000; 2000US-192812P.
XX 29-MAR-2000; 2000US-192916P.
XX 29-MAR-2000; 2000US-192923P.
XX 29-MAR-2000; 2000US-192933P.
XX 29-MAR-2000; 2000US-192945P.
XX (PHNA) PHARMACIA & UPJOHN CO.
XX
XX Vogel G;
XX
XX WPI; 2001-607456/69.
XX P-PSDB; MAU29472.
XX
XX Nucleic acid encoding G protein-coupled receptors, useful for the
XX prevention, diagnosis and treatment of mental disorders.
XX
XX Claim 4; Page 90; 274pp; English.
XX
XX Sequences AAS46819-AA546946 represent cDNA molecules encoding human G
XX protein-coupled receptor (GPCR) polypeptides. The protein and DNA
XX sequences of the invention can be used to identify compounds which bind
XX to GPCR polypeptides and in screening for compounds that modulate GPCR
XX activity. By screening a human subject for the presence of mutations in
XX GPCR DNA, a GPCR-related disorder or a genetic predisposition can be
XX diagnosed. The sequences can also be used for treatment and prevention of
XX mental disorders such as schizophrenia, neurological disorders such as
XX manic depression, metabolic disorders such as obesity, cancer, rheumatoid
XX arthritis, thyroid disorders such as myxoedema, neurodegenerative
XX disorders such as Parkinson's disease, cardiovascular disorders such as
XX atherosclerosis, renal failure, autoimmune disorders, hyperproliferative
XX disorders such as psoriasis and viral infections such as those caused by
XX HIV.
XX
XX Sequence 670 BP; 218 A; 134 C; 143 G; 175 T; 0 other;
SQ
Query Match: 12.2%; Score 61; DP 22; Length: 670;
Best Local Similarity 60.6%; Pred. No. 6.4e-08;
Matches 117; Conservative 0; Mismatches 75; Indels 1; Gaps 1;
QY 237 TGTCTACTTCTCTATCAAGATGTGGAGCTTATTTTCCCATATATTCGACTAGATTGGCT 296
DB :|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
319 TGCAGTGTTCATCAAGAGCTGCATCTTATTTCCCAACCCCAAAATCCA-AGGTGGCT 377
```


PD 07-FEB-2001.
XX 28-JUL-2000; 2000EP-0115126.
XX 29-JUL-1999; 99JP-0248016.
XX 29-AUG-1999; 99JP-0322561.
XX 12-CAN-2000; 2000JP-0118776.
XX 02-MAY-2000; 2000JP-018767.
XX 09-JUN-2000; 2000JP-0241899.
XX (HELI-) HELIX RES INST.
XX Cta T. Tsogai T. Nishikawa T. Hayashi K. Saito K. Yamamoto J.
XX Ishii S. Sugiyama T. Wakamatsu A. Nagai K. Otsuki T.
XX WP1; 2001-318749/34.
XX Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
PT full-length cDNAs.
XX
XX Claim 8; SEQ ID 17036; 2537bp + CD ROM; English.
XX
XX The present invention describes primer sets for synthesizing 5602
XX full-length cDNAs defined in the specification. Where a primer set
XX comprises: (a) an oligo-dr primer and an oligonucleotide complementary
XX to the complementary strand of a polynucleotide which comprises one of
XX the 5602 nucleotide sequences defined in the specification, where the
XX oligonucleotide comprises at least 15 nucleotides; or (b) a combination
XX of an oligonucleotide comprising a sequence complementary to the
XX complementary strand of a polynucleotide which comprises a 5'-end
XX sequence and an oligonucleotide comprising a sequence complementary to a
XX polynucleotide which comprises a 3'-end sequence, where the
XX oligonucleotide comprises at least 15 nucleotides and the combination of
XX the 5'-end sequence/3'-end sequence is selected from those defined in
XX the specification. The primer sets can be used in antisense therapy and
XX in gene therapy. The primers are useful for synthesizing polynucleotides,
XX particularly full-length cDNAs. The primers are also useful for the
XX detection and/or diagnosis of the abnormality of the proteins encoded by
XX the full-length cDNAs. The primers allow obtaining of the full-length
XX cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
XX AAH13631 to AAH18742 represent human cDNA sequences; AAH92446 to
XX AAH35893 represent human amino acid sequences; and AAH13629 to AAH13632
XX represent oligonucleotides, all of which are used in the exemplification
XX of the present invention.
XX
XX Sequence 2477 BP; 697 A; 416 C; 560 G; 804 T; C other;
XX
XX Query Match 11.5%; Score 57.4; DB 12; Length 2477;
XX Best Local Similarity 59.5%; Pred. No. 1.2e-06;
XX Matches 97; Conservative 0; Mismatches 66; Indels 0; Gaps 0;
XX
XX 210 TATATATGCCCCCTTTGCATGTGACCTTTGCTACTTCTTATCAAGATGTGGAGCTTATT 249
XX 944 TATCAACAGCCCTTTGCATGTGACCTTTGCTACTTCTTATCAAGATGTGGAGCTTATT 893
XX 210 TCCCATATATTCATAGATGTGGCTTCTGACTTCTTATCAAGATGTGGAGCTTATT 329
XX 894 TCCCATATATTCATAGATGTGGCTTCTGACTTCTTATCAAGATGTGGAGCTTATT 829
XX 330 ATGACACTGTGGACCTTGGATTTAGTTTCTGAGAGACTTA 172
XX 824 GTACATTTGTGCTGGTTCTGAGCCCAACCTCAGGAGATCCCA 782

Search completed: October 24, 2003, 18:47:15
Job time : 128.251 secs

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ON nucleic - nucleic search, using sw model

Run on: October 24, 2003, 12:34:49 Search time 31.1642 Seconds
(without alignments)
7052.739 Million cell updates/sec

Title: US-09-830-902-1_COPY_4000_4500

Perfect score: 501

Sequence: 1 ccaagtgctggattacag.....gaaatggcagtagtaggc 501

Scoring table: IDENTITY_NJC

Gapop 10.0 , Gapext 1.0

Searched: 569978 seqs, 220591566 residues

Total number of hits satisfying chosen parameters: 1119356

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 2: /cgn2.6/prodata/2/ina/5B_COVB.seq:*
- 3: /cgn2.6/prodata/2/ina/6A_COVB.seq:*
- 4: /cgn2.6/prodata/2/ina/6B_COVB.seq:*
- 5: /cgn2.6/prodata/2/ina/PCUS_COVB.seq:*
- 6: /cgn2.6/prodata/2/ina/backfiles.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	53.6	10.7	1586	3	US-09-173-581-11 Sequence 11, Appl
2	53.6	10.7	1586	3	US-09-173-581-11 Sequence 11, Appl
3	53.6	10.7	1586	4	US-09-387-212-1 Sequence 1, Appl
4	53.6	10.7	1586	4	US-09-387-212-1 Sequence 1, Appl
5	53.6	10.7	2142	4	US-09-948-802-1 Sequence 1, Appl
6	52.2	10.4	2141	4	US-09-441-038-1 Sequence 1, Appl
7	47.8	9.5	537	3	US-08-642-2745-36 Sequence 38, Appl
8	47.8	9.5	537	3	US-08-952-014C-38 Sequence 39, Appl
9	46.6	9.3	485	4	US-09-499-522-5 Sequence 5, Appl
10	45.6	9.1	63588	4	US-09-873-434-3 Sequence 3, Appl
11	45.2	9.0	62450	4	US-09-345-882-1 Sequence 1, Appl
12	45	9.0	3603	4	US-09-705-299-11 Sequence 11, Appl
13	44.4	8.9	43952	4	US-09-735-934A-3 Sequence 3, Appl
14	44.4	8.9	43952	4	US-09-735-934A-3 Sequence 3, Appl
15	44	8.8	17041	3	US-08-076-011-1 Sequence 1, Appl
16	43.9	8.7	3350	3	US-09-110-116-2 Sequence 2, Appl
17	43.8	8.7	17138	3	US-09-613-813-3 Sequence 3, Appl
18	43.8	8.7	17138	4	US-09-803-048-3 Sequence 3, Appl
19	43.8	8.7	64467	3	US-09-803-67-B-3 Sequence 3, Appl
20	43.6	8.7	2373	3	US-09-975-762-45 Sequence 45, Appl
21	43.6	8.7	2373	3	US-09-975-762-45 Sequence 45, Appl
22	43.6	8.7	2373	4	US-09-106-582-45 Sequence 45, Appl
23	43.2	8.6	46718	4	US-09-816-093-3 Sequence 3, Appl
24	43	8.6	41694	4	US-09-536-059-1 Sequence 1, Appl
25	43	8.6	98844	4	US-09-791-211-10 Sequence 10, Appl
26	42.8	8.5	99500	4	US-09-738-094-10 Sequence 10, Appl
27	42.6	8.5	14753	4	US-09-821-736-3 Sequence 3, Appl

C	28	42.2	8.4	35060	3	US-08-914-095-7	Sequence 7, Appl
	29	42.2	8.4	40000	4	US-09-780-049-18	Sequence 18, Appl
	30	42.2	8.4	65942	4	US-09-784-316-3	Sequence 3, Appl
C	31	42.2	8.4	84495	4	US-09-797-906-3	Sequence 3, Appl
	32	42	8.4	776	4	US-09-535-028-37	Sequence 37, Appl
	33	42	8.4	46718	4	US-09-816-093-3	Sequence 3, Appl
C	34	42	8.4	174493	4	US-09-804-47A-3	Sequence 3, Appl
	35	41.8	8.3	36651	4	US-09-738-894A-3	Sequence 3, Appl
	36	41.8	8.3	36651	4	US-09-964-469-3	Sequence 3, Appl
C	37	41.8	8.3	319638	4	US-09-539-333D-1	Sequence 1, Appl
	38	41.8	8.3	319638	4	US-09-679-409-1	Sequence 1, Appl
	39	41.6	8.3	1811	3	US-08-948-253-1	Sequence 1, Appl
C	40	41.6	8.3	36651	4	US-09-738-894A-3	Sequence 3, Appl
	41	41.6	8.3	33651	4	US-09-964-469-3	Sequence 3, Appl
C	42	41.6	8.3	174493	4	US-09-804-47A-3	Sequence 3, Appl
	43	41.6	8.3	4823	2	US-08-457-254-5	Sequence 5, Appl
	44	41.4	8.3	4823	2	US-08-484-257-20	Sequence 20, Appl
	45	41.4	8.3	4823	3	US-09-999-927-5	Sequence 5, Appl

ALIGNMENTS

RESULT 1

US-09-173-581-11
Sequence 11, Application US/09173581A
Patent No. 6033455
GENERAL INFORMATION:
APPLICANT: Bandmar, Olga
APPLICANT: Tang, Y. Tom
APPLICANT: Hillman, Jennifer L.
APPLICANT: Yue, Henry
APPLICANT: Guebler, Karl J.
APPLICANT: Corley, Neil C.
APPLICANT: Gorgone, Gina
APPLICANT: Azimzal, Valda
APPLICANT: Lu, Aina
TITLE OF INVENTION: Protein Kinase Homologs
FILE REFERENCE: PF-0614 US
CURRENT APPLICATION NUMBER: US/09/173,581A
CURRENT FILING DATE: 1998-10-15
NUMBER OF SEQ ID NOS: 18
SOFTWARE: PERL Program
SEQ ID NO 11
LENGTH: 1586
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: 132750
US-09-173-581-11

Query Match 10.7% Score 53.6 DB 3 Length 1586

Best Local Similarity 59.2% Pred No 5.7e-07

Mismatches 103 Conservative 74 Indels 1 Gaps 11

QY	192	CAATTCTCTCTTCATATATATGCCCCCTTTCGAATGACTTGTCTATCTATC	251
DB	1154	CAAAATGTA-CTTCTGGTATGCACACCTCTGGCAATGAAATTCGACCTCCTCCCTC	1213
QY	252	AAAGATGGAGCTATTTTCCCATATATTGCACTAGTTCGCTTCGACTGGCTTGA	311
DB	1214	CATAATGAAGTCTCTTCCCACTATCTGAGTGGCTGGCACTGTGACTGATTGA	1273
QY	312	-CAATGGAATAGTACAAATGCACACTGTGCAACTTGGATTTTAGTTTCGAGAGAACT	370
DB	1274	TCAATGAGATGGGAGAGAGTCACTGTATGCCAGTTCAGGCTAGGTTTCAAGAGGCT	1333
QY	371	TACA 374	
DB	1334	TATA 1337	

RESULT 2

```
US-09-420-915-11
; Sequence 1: Application US/09420915
; Patent No. 6264947
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; APPLICANT: Tang, Y. Tom
; APPLICANT: Hiltner, Jennifer L.
; APPLICANT: Yue, Henry
; APPLICANT: Guegler, Karl J.
; APPLICANT: Corley, Neil C.
; APPLICANT: Gorgone, Gina
; APPLICANT: Azimzai, Valda
; APPLICANT: Lu, Aina
; TITLE OF INVENTION: Protein Kinase Homologs
; FILE REFERENCE: PF-0614 US
; CURRENT APPLICATION NUMBER: US/09/420,915
; CURRENT FILING DATE: 1999-10-20
; EARLIER APPLICATION NUMBER: US 09/173,591
; EARLIER FILING DATE: 1998-10-15
; NUMBER OF SEQ ID NOS: 18
; SOFTWARE: PERL Program
; SEQ ID NO 11
; LENGTH: 1586
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: 132750
US-09-420-915-11

Query Match 10.7%; Score 53.6; DB 3; Length 1586;
Best Local Similarity 59.2%; Pred. No. 6.1e-07;
Matches 109; Conservative 0; Mismatches 74; Indels 1; Gaps 1;

QY 192 CAATTCCTCTATCTCATATATATGCCCCCTTTGCAATGTGACTTGGTACTTCTTATC 251
DB 1154 CAATGTATCTTTCTGGTATCCACACCTCTTGGCAATGGAATTTGACGCTCTCCCTTC 1213
QY 252 AAGATGTGGAGCTTATTTTCCCATATATTTGCACCTAGAGTTGGCTTCTGACTTCTTGA 311
DB 1214 CATTAATGAAGTCTCTTCCCACTTCCACCTCTTGGCAATGGAATTTGACGCTCTCCCTTC 1273
QY 312 -CAATGGAATGTAGTACAAATGACACGTGCAACTTGGATTTAGCTTCGAGAGAACT 370
DB 1274 TCAATGAATGTGAGAGAGTGAATGCTGATCCAGCTTCCAGGCTAGCTTCCAGAGGCGCT 1333
QY 371 TACA 374
DB 1334 TATA 1337

RESULT 3
US-09-387-212-1
; Sequence 1: Application US/09387212A
; Patent No. 6309649
; GENERAL INFORMATION:
; APPLICANT: ROBISON, KEITH E.
; TITLE OF INVENTION: NUCLEIC ACID MOLECULES ENCODING HUMAN KINASE AND
; FILE REFERENCE: MN1-09C
; CURRENT APPLICATION NUMBER: US/09/387,212A
; CURRENT FILING DATE: 1999-08-31
; NUMBER OF SEQ ID NOS: 18
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 1868
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: All occurrences of n indicate any nucleotide
US-09-387-212-1

Query Match 10.7%; Score 53.6; DB 4; Length 1868;
Best Local Similarity 59.2%; Pred. No. 6.1e-07;
Matches 109; Conservative 0; Mismatches 74; Indels 1; Gaps 1;

QY 192 CAATTCCTCTATCTCATATATATGCCCCCTTTGCAATGTGACTTGGTACTTCTTATC 251
DB 1415 CAATGTATCTTTCTGGTATCCACACCTCTTGGCAATGGAATTTGACGCTCTCCCTTC 1474
QY 252 AAGATGTGGAGCTTATTTTCCCATATATTTGCACCTAGAGTTGGCTTCTGACTTCTTGA 311
DB 1476 CATTAATGAAGTCTCTTCCCACTTCCACCTCTTGGCAATGGAATTTGACGCTCTCCCTTC 1534
QY 312 -CAATGGAATGTAGTACAAATGACACGTGCAACTTGGATTTAGCTTCGAGAGAACT 370
DB 1535 TCAATGAATGTGAGAGAGTGAATGCTGATCCAGCTTCCAGGCTAGCTTCCAGAGGCGCT 1594
QY 371 TACA 374
DB 1595 TATA 1598

RESULT 5
US-09-441-039-3
; Sequence 1: Application US/09441039
; Patent No. 6428980
; GENERAL INFORMATION:
; APPLICANT: Luo, Ying
; APPLICANT: Yu, Pei Wen
; APPLICANT: Shen, Mary
; APPLICANT: Huang, Chao Bai
; TITLE OF INVENTION: No. 6428980e1 RFP3 Associated Cell Cycle Proteins, Compositions
```



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; TITLE OF INVENTION: and Methods of Use
; FILE REFERENCE: A68411/DCB/RMS/DAV
; CURRENT APPLICATION NUMBER: US/09/441,039
; CURRENT FILING DATE: 1999-11-16
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 1
; TYPE: DNA
; LENGTH: 2142
; ORGANISM: Homo sapiens
US-09-441-039-1

Query Match      10.7%   Score 53.6; DB 4; Length 2142;
Best Local Similarity 59.2%; Pred. No. 6,48e-07;
Matches 109; Conservative 0; Mismatches 74; Indels 1; Gaps 1;

QY 192 CAATTCTCTATCTCATATATATSCCCCTTTGCAATGTGACTTTGCTACTCTCTATC 251
DB 1698 CAAATGTATCTTTCTGGGTATCCACACACTCTTGGCAATGAAATTTGCAGCTCTCTCTCTG 1759
QY 252 AGATGTGAGGCTTATTTCCCATATATTTGGCACTAGAGTGGCTTCTGACTTGTCTTGA 311
DB 1758 CATAAATGAAGTCTCTTCCCAACCAATTTGAATCTGGGCTGSCACTGTGACTTGATTTGA 1817
QY 312 -CAATGGAATGTAGTACAAATGACACTGTGCCACTTTGGATTTTAGTATTGAGAGAGACT 370
DB 1818 TCAATGAAATGTGGAAGAGAGTACTGTATGCCAATTCGCAAGCCTAGGTTTCAGAGAGGCT 1877
QY 371 TACA 374
DB 1878 TATA 1881

RESULT 6
US-09-441-039-1
; Sequence 1, Application US/09441039
; Patent No. 6428980
; GENERAL INFORMATION:
; APPLICANT: Luo, Ying
; APPLICANT: Yu, Pei Wen
; APPLICANT: Shen, Crac Bai
; APPLICANT: Huang, Crac Bai
; TITLE OF INVENTION: and Methods of Use
; FILE REFERENCE: A68411/DCB/RMS/DAV
; CURRENT APPLICATION NUMBER: US/09/441,039
; CURRENT FILING DATE: 1999-11-16
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 1
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-441-039-1

Query Match      10.4%   Score 52.2; DB 4; Length 2141;
Best Local Similarity 62.2%; Pred. No. 1.7e-04;
Matches 115; Conservative 0; Mismatches 66; Indels 2; Gaps 2;

QY 192 CAATTCTCTATCTCATATATATSCCCCTTTGCAATGTGACTTTGCTACTCTCTATC 250
DB 1696 CAAATGTATCTTTCTGGGTATCCACACACTCTTGGCAATGAAATTTGCAGCTCTCTCTG 1755
QY 251 CAGATGGAGGCTATTTTCCCATATATTTGCCTAGATGGCTTCTGACTTGTCTTGG 310
DB 1756 CCATAAATGAAGTCTCTTCCCAACCAATTTGAATCTGGGCTGSCACTGTGACTTGATTTG 1815
QY 311 A-CAATGGAATGTAGTACAAATGACACTGTGCCACTTTGATTTTAGGTTTCAGAGAGAC 369
DB 1916 ATCAATAGATGTGGAGAGAGTACTGTATGCCAGTTCCAGAGCCTAGGTTTCAGAGAGGCT 1875
QY 370 TTACA 374
DB 1878 TATA 1881

; TITLE OF INVENTION: MUTATED FORMS OF THE ATAXIA-TELANGIECTASIA GENE AND METHOD TO
; SCREEN FOR A PARTIAL A-T PHENOTYPE
; FILE REFERENCE: 229000033
; CURRENT APPLICATION NUMBER: US/08/642,274D
; CURRENT FILING DATE: 1996-05-23
; NUMBER OF SEQ ID NOS: 220
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 38
; TYPE: DNA
; LENGTH: 537
; ORGANISM: Artificial Sequence
; FEATURES: Artificial Sequence
; OTHER INFORMATION: Description of Artificial Sequence: genomic
US-08-642-274D-38

Query Match      9.5%   Score 47.8; DB 3; Length 537;
Best Local Similarity 70.3%; Pred. No. 2.3e-05;
Matches 64; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

QY 1 CCAAAGTGTGGGATTACAGCGGTTACGCACAGCATCCAGCCTTAAGCATGTTAATTAAAG 60
DB 12 CCAAAGTGTGGGATTACAGTCTGTGACCCACCGCCTTAAGCTTAAATTTCTTGA 71
QY 6: TTTTATATTCAGCAAAATGGTTGGAAAT: 91
DB 72 GTACAGAAACACGATTATAGTTTGGAAAT :02

RESULT 8
US-08-952-014C-38
; Sequence 38, Application US/08952014C
; Patent No. 6265158
; GENERAL INFORMATION:
; APPLICANT: Shi,oh, Yosef
; TITLE OF INVENTION: ATAXIA-TELANGIECTASIA GENE AND ITS
; TITLE OF INVENTION: GENOMIC ORGANIZATION
; NUMBER OF SEQUENCES: 91
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kohn & Associates
; STREET: 3050C No. 6265158thwestern Hwy., Suite 413
; CITY: Farmington Hills
; STATE: Michigan
; COUNTRY: U.S.
; ZIP: 48334
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/952,014C
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Kohn, Kenneth
; REGISTRATION NUMBER: 30,995
; REFERENCE/DOCKET NUMBER: 2290, C0028
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 810-539-5050
; TELEFAX: 810-539-5055
; INFORMATION FOR SEQ ID NO: 38:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 537 base pairs
; TYPE: nucleic acid

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STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-952-C14C-38

Query Match          9.5%  Score 47.3; DB 3; Length 537;
Best Local Similarity 70.3%  Pred. No. 2.3e-95;
Matches 64; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

CY 1 CCAAGTGTGGGATTACAGCGCTTAGCCACACATCCAGCCCTTAGCATGTTAATTAG 60
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 12 CCAAGTGTGGGATTACAGCGCTTAGCCACACATCCAGCCCTTAGCATGTTAATTAG 72
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

CY 61 TTTTATAATTCAGCAAAATGGTTGGAAAT 91
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 72 GTACAGAAACACATTAAGTTTGGAAAT 102
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

RESULT 9
US-09-499-522-5/c
; Sequence 5, Application US/09499522
; Patent No. 6479238
; GENERAL INFORMATION:
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Bouqueleret, Lydie
; APPLICANT: Bihain, Bernard
; TITLE OF INVENTION: POLYMORPHIC MARKERS OF THE LSR GENE
; FILE REFERENCE: GENSET.053AUS
; CURRENT APPLICATION NUMBER: US/09/499,522
; EARLIER FILING DATE: 2000-02-10
; EARLIER APPLICATION NUMBER: US 60/119,592
; EARLIER FILING DATE: 1999-02-10
; EARLIER APPLICATION NUMBER: US 60/144,784
; EARLIER FILING DATE: 1998-07-20
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: Patent.pm
; SEQ ID NO 5
; LENGTH: 465
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: allele
; LOCATION: 126
; OTHER INFORMATION: 99-14417-126 : polymorphic base C or T
; FEATURE:
; NAME/KEY: misc_binding
; LOCATION: 103..125
; OTHER INFORMATION: 99-14417-126..151 potential
; FEATURE:
; NAME/KEY: misc_binding
; LOCATION: 127..149
; OTHER INFORMATION: 99-14417-126..152 potential, complement
; FEATURE:
; NAME/KEY: allele
; LOCATION: 334
; OTHER INFORMATION: 99-14417-334 : polymorphic base C or T
; FEATURE:
; NAME/KEY: misc_binding
; LOCATION: 311..333
; OTHER INFORMATION: 99-14417-334..357 potential
; FEATURE:
; NAME/KEY: misc_binding
; LOCATION: 335..357
; OTHER INFORMATION: 99-14417-334..352 potential
; FEATURE:
; NAME/KEY: misc_binding
; LOCATION: 1..21
; OTHER INFORMATION: upstream amplification primer
; FEATURE:
; NAME/KEY: misc_binding
; LOCATION: 447..465
; OTHER INFORMATION: downstream amplification primer, complement
US-09-499-522-5

Query Match          9.3%  Score 46.6; DB 4; Length 463;
Best Local Similarity 59.6%  Pred. No. 5e-05;
Matches 133; Conservative 0; Mismatches 84; Indels 0; Gaps 3;

CY 223 TTGCAATGTGACTTTGCTACT-TCTCTATCAAGATGTGGAGCTTATTTTCCCATATTTG 281
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 434 TTACAAGTGACTTTGGCACTCTTCCCATCAAGGATGGAGTCTGTGTCTCCACGCTTG 375
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

CY 282 CACTAGAGTGGCGCTTCGACTTTCGCTTT-GACAATGGAAATSTAGTACAAATGACACTGTG 340
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 374 AATCGGCTGACCTTGTAAATTTCTTTTGGCCAAATCAACATGGTGAAGTAATGCTGTG 315
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

CY 341 CAACCTTGGATTTTAGGTTTGGAGAGAACTTACACCTTCCACTACACACTCTCTTGGAAAC 400
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 314 TTAGTTTCAAGCCTAGACTTCAAGAAACCTTGGCGCTTCCATGACTCTCTTAGTTGACC 255
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

CY 401 ----CAGATCAATGTAAGAAATCAGGGCTATCTCTGCTAGAG 439
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 254 NTTCCATGACCTTGTGAACATGACCGGCTAGCCTCTCTGGAG 212
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

RESULT 10
US-09-873-424-3
; Sequence 3, Application US/09873404
; Patent No. 6500656
; GENERAL INFORMATION:
; APPLICANT: WEBSTER, Marion et al
; TITLE OF INVENTION: ISOLATED HUMAN KINASE PROTEINS, NUCLEIC
; TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN KINASE PROTEINS, AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CLOC212-CIP
; CURRENT APPLICATION NUMBER: US/09/873,404
; EARLIER FILING DATE: 2001-06-05
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 63583
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1) ..(63586)
; OTHER INFORMATION: n = A,T,C or G
US 09-873-424-3

Query Match          9.1%  Score 45.6; DB 4; Length 63583;
Best Local Similarity 54.9%  Pred. No. 0.00076;
Matches 90; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

CY 1 CCAAAATGCTGGATTACAGCGCTTAGCCACAGCATCCAGCCTTAGCATGTTAATAAG 60
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 417 CCAAAATGCTGGATTACAGCGCTTAGCCACAGCATCCAGCCTTAGCATGTTAATAAG 4230
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

CY 61 TTTTATAATTCAGCAAAATGGTTGGAAATTCGTCTTAAATGAGATGTTTAAGCTGCC 120
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 4231 CATTGCTGTGTATGATATTTACCCCAACTTAGTGGCTTAAGAATAAAGATTTATTGTC 4290
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

CY 121 GTCTGAACATGAGGTAGAGGAATTTCTACACATAATCATTTGTG 164
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 4291 TCAGAGATCCTGTGCATCAAGAA-TTAGGATGACATCAITGAG 4334
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

RESULT 11
US-09-345-882-1/c
; Sequence 1, Application US/09345882
; Patent No. 6395373
; GENERAL INFORMATION:
; APPLICANT: Bouqueleret, Lydie
; TITLE OF INVENTION: A NUCLEIC ACID ENCODING A RETINOBLASTOMA BINDING PROTEIN (RBP2-)
; TITLE OF INVENTION: AND POLYMORPHIC MARKERS ASSOCIATED WITH SAID NUCLEIC ACID.
; FILE REFERENCE: GENSET.031A
; CURRENT APPLICATION NUMBER: US/09/345,882
```

1 CURRENT FILING DATE: 1999-06-30
2 PRIOR APPLICATION NUMBER: US 60/091,375
3 PRIOR FILING DATE: 1998-06-30
4 PRIOR APPLICATION NUMBER: US 60/111,905
5 PRIOR FILING DATE: 1998-12-13
6 NUMBER OF SEQ ID NOS: 140
7 SOFTWARE: Patent-IT
8 SEQ ID NO: 1
9 LENGTH: 162450
10 TYPE: DNA
11 ORGANISM: Homo sapiens
12
13 FEATURE:
14 NAME/KEY: allele
15 LOCATION: 72794
16 OTHER INFORMATION: 5-124-273 : polymorphic base A or G
17 FEATURE:
18 NAME/KEY: allele
19 LOCATION: 95073
20 OTHER INFORMATION: 5-127-261 : polymorphic base A or C
21 FEATURE:
22 NAME/KEY: allele
23 LOCATION: 92842
24 OTHER INFORMATION: 99-1437-325 : polymorphic base A or G
25 FEATURE:
26 NAME/KEY: allele
27 LOCATION: 93714
28 OTHER INFORMATION: 5-128-60 : polymorphic base deletion of GT
29 FEATURE:
30 NAME/KEY: allele
31 LOCATION: 97122
32 OTHER INFORMATION: 99-1442-224 : polymorphic base G or T
33 FEATURE:
34 NAME/KEY: allele
35 LOCATION: 97152
36 OTHER INFORMATION: 5-129-144 : polymorphic base deletion of T
37 FEATURE:
38 NAME/KEY: allele
39 LOCATION: 99096
40 OTHER INFORMATION: 5-130-257 : polymorphic base A or G
41 FEATURE:
42 NAME/KEY: allele
43 LOCATION: 99117
44 OTHER INFORMATION: 5-130-276 : polymorphic base A or G
45 FEATURE:
46 NAME/KEY: allele
47 LOCATION: 103805
48 OTHER INFORMATION: 5-131-395 : polymorphic base A or C
49 FEATURE:
50 NAME/KEY: allele
51 LOCATION: 106940
52 OTHER INFORMATION: 5-133-375 : polymorphic base insertion of A
53 FEATURE:
54 NAME/KEY: allele
55 LOCATION: 108136
56 OTHER INFORMATION: 5-135-155 : polymorphic base insertion of A
57 FEATURE:
58 NAME/KEY: allele
59 LOCATION: 108149
60 OTHER INFORMATION: 5-135-198 : polymorphic base insertion of GTT
61 FEATURE:
62 NAME/KEY: allele
63 LOCATION: 108308
64 OTHER INFORMATION: 5-135-357 : polymorphic base A or G
65 FEATURE:
66 NAME/KEY: allele
67 LOCATION: 108471
68 OTHER INFORMATION: 5-136-174 : polymorphic base C or T
69 FEATURE:
70 NAME/KEY: allele
71 LOCATION: 134334
72 OTHER INFORMATION: 5-140-123 : polymorphic base C or T
73 FEATURE:
74 NAME/KEY: allele
75 LOCATION: 134362
76 OTHER INFORMATION: 5-140-348 : polymorphic base insertion of A
77 FEATURE:
78 NAME/KEY: allele
79 LOCATION: 134374
80 OTHER INFORMATION: 5-140-361 : polymorphic base insertion of CA
81 FEATURE:
82 NAME/KEY: allele
83 LOCATION: 146328
84 OTHER INFORMATION: 5-143-84 : polymorphic base A or G
85 FEATURE:
86 NAME/KEY: allele
87 LOCATION: 146345
88 OTHER INFORMATION: 5-143-101 : polymorphic base A or C
89 FEATURE:
90 NAME/KEY: allele
91 LOCATION: 150323
92 OTHER INFORMATION: 5-145-24 : polymorphic base A or G
93 FEATURE:
94 NAME/KEY: allele
95 LOCATION: 160031
96 OTHER INFORMATION: 5-148-352 : polymorphic base G or T
97 FEATURE:
98 NAME/KEY: allele
99 LOCATION: 72771..72817
100 OTHER INFORMATION: polymorphic fragment 5-124-273 SEQ ID30
101 FEATURE:
102 NAME/KEY: allele
103 LOCATION: 72771..72817
104 OTHER INFORMATION: polymorphic fragment 5-124-273 SEQ ID51
105 FEATURE:
106 NAME/KEY: allele
107 LOCATION: 88050..88096
108 OTHER INFORMATION: polymorphic fragment 5-127-261 SEQ ID32
109 FEATURE:
110 NAME/KEY: allele
111 LOCATION: 88050..88096
112 OTHER INFORMATION: polymorphic fragment 5-127-261 SEQ ID52
113 FEATURE:
114 NAME/KEY: allele
115 LOCATION: 93819..93865
116 OTHER INFORMATION: complement polymorphic fragment 99-1437-325 SEQ ID49
117 FEATURE:
118 NAME/KEY: allele
119 LOCATION: 93819..93865
120 OTHER INFORMATION: complement polymorphic fragment 99-1437-325 SEQ ID70
121 FEATURE:
122 NAME/KEY: allele
123 LOCATION: 93690..93736
124 OTHER INFORMATION: polymorphic fragment 5-128-60 SEQ ID32
125 FEATURE:
126 NAME/KEY: allele
127 LOCATION: 93690..93736
128 OTHER INFORMATION: polymorphic fragment 5-128-60 SEQ ID53
129 FEATURE:
130 NAME/KEY: allele
131 LOCATION: 97099..97145
132 OTHER INFORMATION: polymorphic fragment 99-1442-224 SEQ ID50
133 FEATURE:
134 NAME/KEY: allele
135 LOCATION: 97099..97145
136 OTHER INFORMATION: polymorphic fragment 99-1442-224 SEQ ID71
137 FEATURE:
138 NAME/KEY: allele
139 LOCATION: 97130..97177
140 OTHER INFORMATION: polymorphic fragment 5-129-144 SEQ ID33
141 FEATURE:
142 NAME/KEY: allele
143 LOCATION: 97130..97177
144 OTHER INFORMATION: polymorphic fragment 5-129-144 SEQ ID54
145 FEATURE:
146 NAME/KEY: allele
147 LOCATION: 99375..99421


```

, CURRENT FILING DATE: 2002-02-01
, NUMBER OF SEQ ID NOS: 4
, SOFTWARE: FastSeq for Windows Ver
, SEQ ID NO 1
, LENGTH: 4395
, TYPE: DNA
, ORGANISM: Homo sapiens
US-0-060-332-3

```

Query Match: 8.9%; Score 44.4; 25.4; Length 43250;
Best Local Similarity 77.1%; Pred. No. 0.0015;
Matches 54; Conservative 37; Mismatches 16; Indels 37

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GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.
OM nucleic - nucleic search, using sw model
Run on: October 24, 2003, 15:28:54 Search time 133.415 Seconds
without alignments: 10586.662 Million cells updates/sec
Title: US-09-830-902-1_COPY_4000_4500
Perfect score: 50
Sequence: 1 CCAAGAGTGGGATTACAG.....TAAATGAGGCTAGTAGGTT 561
Scoring table: IDENTITY_NUC
Gapop 15.0, Gapext 1.0
Searched: 1792395 seqs, 134590451 residues
Total number of hits satisfying chosen parameters: 3584730
Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 53
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications, NA:
1: /cgn2_6/prodata/2/pubpna/US07_PUBCOMB.seq:
2: /cgn2_6/prodata/2/pubpna/PCT_NEW_PUB.seq:
3: /cgn2_6/prodata/2/pubpna/US06_NEW_PUB.seq:
4: /cgn2_6/prodata/2/pubpna/US06_PUBCOMB.seq:
5: /cgn2_6/prodata/2/pubpna/US07_NEW_PUB.seq:
6: /cgn2_6/prodata/2/pubpna/PCTUS_PUBCOMB.seq:
7: /cgn2_6/prodata/2/pubpna/US08_NEW_PUB.seq:
8: /cgn2_6/prodata/2/pubpna/US08_PUBCOMB.seq:
9: /cgn2_6/prodata/2/pubpna/US09_PUBCOMB.seq:
10: /cgn2_6/prodata/2/pubpna/US09_PUBCOMB.seq:
11: /cgn2_6/prodata/2/pubpna/US09_PUBCOMB.seq:
12: /cgn2_6/prodata/2/pubpna/US09_NEW_PUB.seq:
13: /cgn2_6/prodata/2/pubpna/US10_PUBCOMB.seq:
14: /cgn2_6/prodata/2/pubpna/US10_PUBCOMB.seq:
15: /cgn2_6/prodata/2/pubpna/US10_NEW_PUB.seq:
16: /cgn2_6/prodata/2/pubpna/US10_PUBCOMB.seq:
17: /cgn2_6/prodata/2/pubpna/US10_PUBCOMB.seq:
Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES				Query		Match		Length		DB		Description	
Result No.	Score	Match	Length	DB	ID	Score	Match	Length	DB	ID	Score	Match	Description
1	68.8	13.7	756	13	US-10-027-632-18943	Sequence 18943, A							
2	68.8	13.7	756	13	US-10-027-632-149822	Sequence 149822, A							
3	64.6	12.9	586	12	US-10-029-386-12215	Sequence 12215, A							
C 4	64.6	12.9	5239	12	US-10-311-455-246	Sequence 246, App							
C 5	64.6	12.9	3673778	12	US-10-312-841-2	Sequence 2, Appl							
C 6	61	12.2	670	9	US-09-811-264-93	Sequence 53, Appl							
7	60.6	12.1	624	13	US-10-027-632-55107	Sequence 55107, A							
8	60.6	12.1	624	13	US-10-027-632-31174	Sequence 31174, A							
9	60.6	12.1	624	13	US-10-027-632-908	Sequence 908, App							
10	60.4	12.1	5239	12	US-10-311-455-245	Sequence 245, App							
11	60.4	12.1	3673778	12	US-10-312-841-1	Sequence 1, Appl							
C 12	60.2	12.0	651	13	US-10-040-739-925	Sequence 925, App							
C 13	59.4	11.9	594	13	US-10-027-632-67207	Sequence 67207, A							
C 14	56.6	11.3	646	13	US-10-027-632-16407	Sequence 16407, A							
C 15	55	11.0	546	13	US-10-027-632-160975	Sequence 160975, A							
C 16	55	11.0	546	13	US-10-027-632-160976	Sequence 160976, A							

C 17	55	11.0	546	13	US-10-027-632-160977	Sequence 160977, A
C 18	54.6	10.9	442	13	US-10-027-632-318161	Sequence 318161, A
C 19	54.6	10.9	450	13	US-10-027-632-93145	Sequence 93145, A
C 20	53.8	10.7	590	13	US-10-027-632-85250	Sequence 85250, A
C 21	53.8	10.7	590	13	US-10-027-632-314433	Sequence 314433, A
C 22	53.8	10.7	604	13	US-10-027-632-105017	Sequence 105017, A
C 23	53.8	10.7	624	13	US-10-027-632-214356	Sequence 214356, A
C 24	53.8	10.7	726	13	US-10-027-632-13713	Sequence 13713, A
C 25	53.6	10.7	586	9	US-09-870-982-1	Sequence 1, Appl
C 26	53.6	10.7	242	14	US-10-165-696-3	Sequence 3, Appl
C 27	53.6	10.7	584	13	US-10-027-632-46655	Sequence 46655, A
C 28	53.4	10.7	584	13	US-10-027-632-295733	Sequence 295733, A
C 29	53.4	10.7	584	13	US-10-027-632-270803	Sequence 270803, A
C 30	53.2	10.6	716	13	US-10-027-632-270824	Sequence 270824, A
C 31	53.2	10.6	716	13	US-10-027-632-174127	Sequence 174127, A
C 32	53.2	10.6	752	13	US-10-027-632-138083	Sequence 138083, A
C 33	53.2	10.6	815	13	US-10-027-632-227143	Sequence 227143, A
C 34	52.8	10.5	556	13	US-09-813-320-3	Sequence 3, Appl
C 35	52.6	10.4	214	14	US-10-165-696-1	Sequence 1, Appl
C 36	52.2	10.4	857	13	US-10-027-632-167271	Sequence 167271, A
C 37	52	10.4	857	13	US-10-027-632-167272	Sequence 167272, A
C 38	52	10.4	857	13	US-10-027-632-167273	Sequence 167273, A
C 39	52	10.4	857	13	US-10-027-632-167274	Sequence 167274, A
C 40	52	10.4	857	13	US-09-764-855-163	Sequence 163, App
C 41	51.6	10.3	1434	14	US-10-072-349-163	Sequence 163, App
C 42	51.6	10.3	1434	14	US-10-027-632-33733	Sequence 33733, A
C 43	50.6	10.1	702	13	US-10-027-632-51013	Sequence 51013, A
C 44	50.4	10.1	522	13	US-10-027-632-85803	Sequence 85803, A
C 45	50.4	10.1	522	13	US-10-027-632-85803	Sequence 85803, A

ALIGNMENTS

RESULT :
US-10-027-632-18943
Sequence 18943, Application US:10027532
GENERAL INFORMATION:
APPLICANT: Wang, David G.
TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
TITLE OF INVENTION: Polymorphisms in the Human Genome
FILE REFERENCE: 108827.129
CURRENT APPLICATION NUMBER: US/10/027.632
PRIOR FILING DATE: 2002-04-30
PRIOR APPLICATION NUMBER: US 60/218,006
PRIOR FILING DATE: 2000-07-12
PRIOR APPLICATION NUMBER: US 60/198,676
PRIOR FILING DATE: 2000-04-20
PRIOR APPLICATION NUMBER: US 60/193,483
PRIOR FILING DATE: 2000-03-29
PRIOR APPLICATION NUMBER: US 60/185,218
PRIOR FILING DATE: 2000-02-24
PRIOR APPLICATION NUMBER: US 60/167,363
PRIOR FILING DATE: 1999-11-23
PRIOR APPLICATION NUMBER: US 60/156,358
PRIOR FILING DATE: 1999-09-28
PRIOR APPLICATION NUMBER: US 60/146,002
NUMBER OF SEQ ID NOS: 325720
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 18943
LENGTH: 756
TYPE: DNA
ORGANISM: Human
US-10-027-632-18943

Query Match 13.7%; Score 69.8; DB 13; Length 756;
Best Local Similarity 62.5%; Pred. No. 1e-09;
Matches 140; Conservative 0; Mismatches 82; Indels 2; Gaps 2;
CY 156 ATCATTTGGCTAAATTAATTCCTGCAGACAAATTCCTATATATATA 215
DB 169 ATCATTTGGCTAAATTCCTGCAGACAAATTCCTATATATATA 168


```
/ APPLICANT: Wang, David G.
/ TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
/ FILE OF INVENTION: Polymorphisms in the Human Genome
/ FILE REFERENCE: 108827.129
/ CURRENT FILING DATE: 2002-04-30
/ PRIOR APPLICATION NUMBER: US 60/218,006
/ PRIOR FILING DATE: 2000-07-12
/ PRIOR APPLICATION NUMBER: US 60/198,676
/ PRIOR FILING DATE: 2000-04-20
/ PRIOR APPLICATION NUMBER: US 60/193,483
/ PRIOR FILING DATE: 2000-03-29
/ PRIOR APPLICATION NUMBER: US 60/185,218
/ PRIOR FILING DATE: 2000-02-24
/ PRIOR APPLICATION NUMBER: US 60/167,363
/ PRIOR FILING DATE: 1999-11-23
/ PRIOR APPLICATION NUMBER: US 60/156,358
/ PRIOR FILING DATE: 1999-09-28
/ PRIOR APPLICATION NUMBER: US 60/146,002
/ PRIOR FILING DATE: 1999-08-09
/ NUMBER OF SEQ ID NOS: 325720
/ SOFTWARE: FastSEQ for Windows Version 4.0
/ SEQ ID NO 55:07
/ LENGTH: 624
/ TYPE: DNA
/ ORGANISM: Human
/ US-10-027-632-55167

Query Match      12.1%  Score 60.6;  DB 13;  Length 624;
Best Local Similarity 61.7%;  Pred. No. 2.3e-07;
Matches 113;  Conservative 0;  Mismatches 69;  Indels 1;  Gaps 1;

QY 179 AAGATGGCCACACAAATTCCTCTATCCCTCATATATATGCCCCCTTGCATATGTGACTTTG 238
DB 442 AAAAATACCTCCCATGTCTCCACCCCTTCTCTGGTATACACATACATATATATGGCTTTG 501
QY 239 CTACTTCTCTATCAAGATGGAGCTTATTTCCCATATATTTCCCATATATTCGACTAGAGTTGGCTTC 298
DB 502 CTGCTCTTCTCATCAAGATGGAGTGGATCGACTTACTTAATCTGATGTGAGCTGGCTTG 561
QY 299 TGACTTGCCTTGA-CAATGGAATGTAGTACAAATGACACTGTGCAACTTTGGATTTTAGG 357
DB 562 TAACTTGCCTTGGACCAATGGAATGTGCCAGAGCGACCTTGTSCCAATTCGTAGTCTAAG 621
QY 358 TTT 360
DB 622 CTT 624

RESULT 8
US-10-027-632-311174
/ Sequence 311174: Application US/10027632
/ GENERAL INFORMATION:
/ APPLICANT: Wang, David G.
/ TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
/ FILE OF INVENTION: Polymorphisms in the Human Genome
/ FILE REFERENCE: 108827.129
/ CURRENT APPLICATION NUMBER: US/10/027.632
/ CURRENT FILING DATE: 2002-04-30
/ PRIOR APPLICATION NUMBER: US 60/218,006
/ PRIOR FILING DATE: 2000-07-12
/ PRIOR APPLICATION NUMBER: US 60/198,676
/ PRIOR FILING DATE: 2000-04-20
/ PRIOR APPLICATION NUMBER: US 60/193,483
/ PRIOR FILING DATE: 2000-03-29
/ PRIOR APPLICATION NUMBER: US 60/185,218
/ PRIOR FILING DATE: 2000-02-24
/ PRIOR APPLICATION NUMBER: US 60/167,363
/ PRIOR FILING DATE: 1999-11-23
/ PRIOR APPLICATION NUMBER: US 60/156,358
/ PRIOR FILING DATE: 1999-09-28
/ PRIOR APPLICATION NUMBER: US 60/146,002
/ NUMBER OF SEQ ID NOS: 325720
/ SOFTWARE: FastSEQ for Windows Version 4.0
/ SEQ ID NO 908
/ LENGTH: 627
/ TYPE: DNA
/ ORGANISM: Human
/ US-10-027-632-908

Query Match      12.1%  Score 60.6;  DB 13;  Length 627;
Best Local Similarity 61.7%;  Pred. No. 2.3e-07;
Matches 113;  Conservative 0;  Mismatches 69;  Indels 1;  Gaps 1;

QY 179 AAGATGGCCACACAAATTCCTCTATCCCTCATATATATGCCCCCTTGCATATGTGACTTTG 238
DB 442 AAAAATACCTCCCATGTCTCCACCCCTTCTCTGGTATACACATACATATATATGGCTTTG 503
QY 239 CTACTTCTCTATCAAGATGGAGCTTATTTCCCATATATTCGACTAGAGTTGGCTTC 298
DB 504 CTGCTCTTCTCATCAAGATGGAGTGGATCGACTTACTTAATCTGATGTGAGCTGGCTTG 563
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/ NUMBER OF SEQ ID NOS: 325720
/ SOFTWARE: FastSEQ for Windows Version 4.0
/ SEQ ID NO 311174
/ LENGTH: 624
/ TYPE: DNA
/ ORGANISM: Human
/ US-10-027-632-311174

Query Match      12.1%  Score 60.6;  DB 13;  Length 624;
Best Local Similarity 61.7%;  Pred. No. 2.3e-07;
Matches 113;  Conservative 0;  Mismatches 69;  Indels 1;  Gaps 1;

QY 179 AAGATGGCCACACAAATTCCTCTATCCCTCATATATATGCCCCCTTGCATATGTGACTTTG 238
DB 442 AAAAATACCTCCCATGTCTCCACCCCTTCTCTGGTATACACATACATATATATGGCTTTG 501
QY 239 CTACTTCTCTATCAAGATGGAGCTTATTTCCCATATATTCGACTAGAGTTGGCTTC 298
DB 502 CTGCTCTTCTCATCAAGATGGAGTGGATCGACTTACTTAATCTGATGTGAGCTGGCTTG 561
QY 299 TGACTTGCCTTGA-CAATGGAATGTAGTACAAATGACACTGTGCAACTTTGGATTTTAGG 357
DB 562 TAACTTGCCTTGGACCAATGGAATGTGCCAGAGCGACCTTGTSCCAATTCGTAGTCTAAG 621
QY 358 TTT 360
DB 622 CTT 624

RESULT 9
US-10-027-632-908
/ Sequence 908: Application US/10027632
/ GENERAL INFORMATION:
/ APPLICANT: Wang, David G.
/ TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
/ FILE OF INVENTION: Polymorphisms in the Human Genome
/ FILE REFERENCE: 108827.129
/ CURRENT APPLICATION NUMBER: US/10/027.632
/ CURRENT FILING DATE: 2002-04-30
/ PRIOR APPLICATION NUMBER: US 60/218,006
/ PRIOR FILING DATE: 2000-07-12
/ PRIOR APPLICATION NUMBER: US 60/198,676
/ PRIOR FILING DATE: 2000-04-20
/ PRIOR APPLICATION NUMBER: US 60/193,483
/ PRIOR FILING DATE: 2000-03-29
/ PRIOR APPLICATION NUMBER: US 60/185,218
/ PRIOR FILING DATE: 2000-02-24
/ PRIOR APPLICATION NUMBER: US 60/167,363
/ PRIOR FILING DATE: 1999-11-23
/ PRIOR APPLICATION NUMBER: US 60/156,358
/ PRIOR FILING DATE: 1999-09-28
/ PRIOR APPLICATION NUMBER: US 60/146,002
/ NUMBER OF SEQ ID NOS: 325720
/ SOFTWARE: FastSEQ for Windows Version 4.0
/ SEQ ID NO 908
/ LENGTH: 627
/ TYPE: DNA
/ ORGANISM: Human
/ US-10-027-632-908

Query Match      12.1%  Score 60.6;  DB 13;  Length 627;
Best Local Similarity 61.7%;  Pred. No. 2.3e-07;
Matches 113;  Conservative 0;  Mismatches 69;  Indels 1;  Gaps 1;

QY 179 AAGATGGCCACACAAATTCCTCTATCCCTCATATATATGCCCCCTTGCATATGTGACTTTG 238
DB 444 AAAAATACCTCCCATGTCTCCACCCCTTCTCTGGTATACACATACATATATATGGCTTTG 503
QY 239 CTACTTCTCTATCAAGATGGAGCTTATTTCCCATATATTCGACTAGAGTTGGCTTC 298
DB 504 CTGCTCTTCTCATCAAGATGGAGTGGATCGACTTACTTAATCTGATGTGAGCTGGCTTG 563
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/ CLASSIFICATION: <Unknown>
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: 09/036,522
/ FILING DATE: 03-JUN-1998
/ ATTORNEY/AGENT INFORMATION:
/ NAME: BROWN, SCOTT A.
/ REGISTRATION NUMBER: 12,724
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: (617) 498-8224
/ TELEFAX: (617) 876-5951
/ INFORMATION FOR SEQ ID NO: 925:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 65 base pairs
/ TYPE: nucleic acid
/ STRANDEDNESS: double
/ TOPOLOGY: linear
/ MOLECULE TYPE: cDNA
/ SEQUENCE DESCRIPTION: SEQ ID NO: 925:
US-10-040-739-925

Query Match      12.0%; Score 60.2; DB 13; Length 651;
Best Local Similarity 61.6%; Pred. No. 3.1e-07;
Matches 146; Conservative 0; Mismatches 88; Indels 3; Gaps 3;

QY 192 ATGCCACACAAATTCCTCTCTCCATATATATATGCCCCCTTTGGAAATGAGCTTTGCTA 241
DB 368 AAGGCCACAAATGAGTAATCTCTCCATGTCAGACTCTTTGGAAATGAGCTTTGCTA 309
QY 242 CTCTCTATCAAGATGGAGCTTAATTTCCCATATATTCACATAGATGGCTTTCTGA 301
DB 308 CTCTCTC-ACGAGMAATGAGCTATTTTCCATCTTTTGAATCTGGCTGGCTTGTGA 250
QY 302 CTCTCTTTGAC-AATGGAATCTAGTACAAATGACATGTGCACTTTTGGATTTTGGCTT 359
DB 249 CTCTCTTTGACTACACAAAGTGGCAGACATCATCTGTATGAGTCTCTGGCATAGGGG 390
QY 360 TCGAGAGAACTTACACTTCCACTCCACACTCTCTTGAACACACAGATGCACGTGAAG 416
DB 189 TTAGAGGATTTGCAATCTTACTCTTTCTTTTGGACCTTAAGCCATGATGA-G 433
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RESULT 13
US-10-027-632-67207/c
; Sequence 67207, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO: 67207
; LENGTH: 594
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-67207
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/ Query Match      11.9%; Score 59.4; DB 13; Length 594;
/ Best Local Similarity 63.3%; Pred. No. 5.1e-07;
/ Matches 107; Conservative 0; Mismatches 61; Indels 1; Gaps 1;

QY 157 TCATTGTGCTAATTAATTTGCAAGATGGCCACAAATTCCTCTATCCTCATATATAT 216
DB 324 TTAGTGTGCTCAATGTTTSCAGAGTACACTTCTGCTATCTCTGTGTGCTGT 263
QY 217 GCGCTTTGCAATGAGCTTTGCTACTTCTCTATCAAGATGTGAGCTTATTTCCCAT 295
DB 264 GCGCTTTGCGGGTGACTTTGCGAGTCTCCCATGGTACAGTGGCATCTGTCTTCCCTT 205
QY 276 AATTGCACTAGATGGCTTCTGACTTCTGCTTTGCAATGCAATGCAATGTAG 324
DB 204 CATCTGTGCTGGCTAGGCTCTGACTTCTTTTGAATCAAAACTTG 156
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RESULT 14
US-10-027-632-16407/c
; Sequence 16407, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/219,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO: 16407
; LENGTH: 646
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-16407
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/ Query Match      11.3%; Score 54.6; DB 13; Length 646;
/ Best Local Similarity 69.4%; Pred. No. 3.5e-06;
/ Matches 77; Conservative 0; Mismatches 34; Indels 2; Gaps 2;

QY 393 TCACACTCTCTTGGAAACGAGATGCAATGTAAGAAGTCAGGGCTATCTCTGTAGACACA 442
DB 451 TCTCTCTCTCTTGGAGGCGAGATGCCATTTGAAAAGACTGGCTCATCTCTGTGAAGACA 392
QY 443 TATGTCCTCCAGCTAATAGCCCAATCACTCTCTGAACATATGAATGAGCTA 493
DB 391 TGTGGCATAGATTATAGCCCAACCAATTCCTCATCTCTGTGAGTGAGGCCA 341
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RESULT 15
US-10-027-632-160975/c
; Sequence 160975, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
```

```

; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,219
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO: 160975
; LENGTH: 546
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-160975

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Query Match      11.83; Score 55; DB 13; Length 546;
Best Local Similarity 62.18; Freq. No. 9.3e-06;
Matches 100; Conservative 3; Mismatches 53; Indels 5; Gaps 17
QY      206 CTGATATATATGCGCCCTTGCAGTGGAGCTTGTCTGCTATATCATCAAGATGGTGGAGCTT 265
DB      283 CCTGATCAATACCCCTTGCATGTGATTTTAAATGCTGATCAATATATATATATATAT 229
QY      266 ATTTCCATATATGGCACTAGAGTTGGCTTCTGATTTCTTTTCAATGGATGTAGT 125
DB      228 ATTTCTACCCCTTGTGANTTGGCTGGCTGCTTCTGCTTCTTTTGGCAATAAATGCAAT 169
QY      326 ACAATGACACTGGCACTTGGATTTTAGGTTTCAGAG 166
DB      168 AGAAGTGATAGTGAACCAAGTCCCAAGTCTGGGCTTAAACAG 123

```

Search completed: October 24, 2003, 18:34:26
Job time : 134.415 secs

GenCore version 5.1.6
Copyright: (c) 1993 - 2003 CompuGen Ltd.

DOM nucleic - nucleic search, using sw model

Run on: October 24, 2003, 18:12:56 : Search time 1546.7 Seconds
(without alignments)
1:633.309 Million cell updates/sec

Title: US-09-830-902-1_COPY_4000_4500

Perfect score: 501

Sequence: 1 ccaaaagtcctgggattacag.....tgaatgaggtacgtatgagcc 501

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 22791332 seqs, 1215223856 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Database :

EST:*

1: em_estba:*

2: em_esthum:*

3: em_estinv:*

4: em_estmni:*

5: em_estov:*

6: em_estpl:*

7: em_estro:*

8: em_esth:*

9: gb_est1:*

10: gb_est2:*

11: gb_est3:*

12: gb_est4:*

13: gb_est5:*

14: gb_est6:*

15: em_estfun:*

16: em_estom:*

17: em_gss_hum:*

18: em_gss_inv:*

19: em_gss_p.n:*

20: em_gss_vit:*

21: em_gss_fun:*

22: em_gss_mni:*

23: em_gss_mus:*

24: em_gss_pro:*

25: em_gss_rdi:*

26: em_gss_pig:*

27: em_gss_val:*

28: gb_gss1:*

29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	ID	Description
C 1	85	17.0	746	29	AG030296 Pan troglodytes
C 2	77	15.4	670	29	BZ610078 HAPD231F
C 3	76.6	15.3	657	14	CB215565 NISC mp27
C 4	73.8	14.7	442	28	B85946 Ctr-HSP-22

5	71	14.2	655	29	AG046887
6	70.6	14.1	608	28	AO35-210
7	67.9	13.5	555	28	AO614358 HS 5126-B
8	67.4	13.5	722	13	EX103739
9	66.4	13.3	586	9	AL733889
10	65.6	13.1	477	9	A104-146
11	65	13.0	474	10	BF329327
12	64.8	12.9	401	28	AO80797
13	64.5	12.9	1335	11	AK046951
14	63.9	12.7	865	20	CC142611
15	63.6	12.7	497	13	BX116065
16	63.4	12.7	528	13	BQ56781
17	63	12.6	726	28	AO389497
18	62.2	12.4	492	28	B33328
19	62	12.4	468	28	AO3394-10
20	62	12.4	540	28	AO175020
21	61.4	12.3	526	28	AO175021
22	59.6	11.9	457	28	AO769797
23	59.2	11.8	373	28	B83258
24	59.2	11.8	498	28	AO1411-4
25	59	11.8	502	2	HSY073396
26	59	11.8	523	10	BF435562
27	58.6	11.7	510	14	CB270506
28	57.6	11.3	1201	9	AJ579387
29	56.8	11.1	680	13	BU685340
30	55	11.0	511	28	AO894910
31	54.2	10.8	421	28	AO194975
32	53.6	10.7	453	9	A1334217
33	53.6	10.7	453	9	AA586672
34	53.6	10.7	459	14	CB069503
35	53.6	10.7	461	28	AO033949
36	53.6	10.7	469	9	A1573185
37	53.6	10.7	506	9	AA706818
38	53.6	10.7	524	13	BU076998
39	53.6	10.7	556	14	CD369377
40	53.6	10.7	622	14	CA427318
41	53.6	10.7	633	14	CB528830
42	53.6	10.7	666	14	CD370383
43	53.6	10.7	670	13	BC575071
44	53.6	10.7	697	13	BX166136
45	53.6	10.7	956	13	BX352121

ALIGNMENTS

RESULT 1
AG030296/c
LOCUS
DEFINITION Pan troglodytes DNA, clone: PTB-002K06.R, genomic survey sequence.
ACCESSION AG030296
VERSION AG030296.1 GI:16557169
KEYWORDS GSS.
SOURCE Pan troglodytes (chimpanzee);
ORGANISM
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.

AG030296 726 bp DNA linear GSS 01-NOV-2001
Pan troglodytes DNA, clone: PTB-002K06.R, genomic survey sequence.

REFERENCE
AUTHORS
Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T.,
Tokoki, Y., Watanabe, H. and Sakaki, Y.
TITLE
RAC end sequences of Library PTB
JOURNAL
Unpublished
2 (bases 1 to 726)

REFERENCE
AUTHORS
Fujiyama, A., Hattori, M., Toyoda, A., Taylor, T. D., Yada, T.,
Tokoki, Y., Watanabe, H. and Sakaki, Y.
TITLE
Direct Submission
JOURNAL
Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0345, Japan
E-mail: chimbesgsc.riken.go.jp, URL: http://hgp.gsc.riken.go.jp/
Tel: 81-45-503-9111, Fax: 81-45-503-9173;
Clones are derived from the chimpanzee BAC library PTB This BAC end
was generated during the R&D process and may have higher chance of


```

/lab_host="DH10B (TI-resistant)"
/clone_lib="NICHD HS Ut1"
/clone="Organ: uterus; Vector: PCMV-SP616.1 ccds (ResGen,
Invitrogen Corporation); Site_1: Not1; Site_2: EcoRV;
Cloned unidirectionally from microquantity amounts of mRNA
from normal endometrial tissue (late proliferative phase,
cycle day 13). Average insert size 1.9 kb. Library
constructed by ResGen (Invitrogen Corporation)."
BASE COUNT      196 a 156 c 125 g 182 t
ORIGIN
Query Match      15.3%; Score 76.6; DP 14; Length 657;
Best Local Similarity 62.3%; Pred. No. 0.0073;
Matches 154; Conservative 0; Mismatches 89; Indels 4; Gaps 2;
QY 196 TCTCTGATCCCATATATATGCGCCCTTTGGCATGTGACTTGGCTACTTCTATCAAGA 255
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 5 TCGACCTTCCCTGGATACACACTATATATATATATATATATATATATATATATATAT 64
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 256 TGTGAGGCTTATTTCCCATATATATGACATAGATGTGCTTTGACTTGGCTTGA-GAA 314
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 65 ATGGAAGGACTTACCTAATGCTTGA-GTGAAGCTGGCTTGTATGCTTGTGACCA 124
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 315 TGGATGTAGTACAAATGACCTGTGCACTTTGATTTAGCTTTCGAGAGAACTTACA 374
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 125 TGGAAATGTCCAGAGCGACCTGTGCCAATTTGAGTCTAAGCTCAAGAGCGCTTGTG 184
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 375 CTTTCA--CTCACACTCTCTGGAAACGACATGCAATGTAAGAAATGAGGCTATCC 431
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 185 TGTTCACACTCTCCCTTTAGAGCCCTGACAGATACCATGTCATGAGCCCGGTAGCC 244
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 432 TGTCTAGA 438
DB |||||
QY 245 TGTCTAGA 251
DB |||||

B80948      442 bp      DNA      linear      GSS 25-JUN-1998
CIT-HSP-2025N4.TPB CIT-HSP Homo sapiens genomic clone 2025N4,
genomic survey sequence.
B80948.1 GI:2867971
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 442)
Adams,M.D., Rounsley,S.D., Zhao,S., Field,C.E., Bass,S., Linber,K.,
Golden,K., Betry,K., Granger,D., Suh,E., White,C., Shizuya,H.,
Simon,M. and Venter,J.C
Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: mdadams@tigr.org
Clones are available from Research Genetics (info@resgen.com). BAC
end search page:
http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.htm
Seq primer: M13-21
Class: BAC ends.
Location/Qualifiers
1..442
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="GDB:7047102"

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/db_xref="taxon:9606"
/clone="2025N4"
/sex="Male"
/cell_type="Sperm"
/clone_lib="CIT-HSP"
/notes="Vector: pBeLoBAC11; Site_1: HindIII; Site_2:
HindIII"
BASE COUNT      133 a 111 c 93 g 105 t
ORIGIN
Query Match      14.7%; Score 73.8; DP 28; Length 442;
Best Local Similarity 60.5%; Pred. No. 0.027;
Matches 147; Conservative 0; Mismatches 97; Indels 5; Gaps 1;
QY 154 TAATCTTTGCTCAAAATTAAGTTCGAAGATGGCAGCAAGATTCCTCTATCTCATATA 213
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 319 TAGAAGTGTGGTAGTGTCTTGCAGAGATGACCAATCAATTCCTTTATCTCTCG---- 284
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 214 TATGCCCCCTTGGCAATGTGACTTTGCTTACTTCTATCAAGAATGGAGGCTTATTTCCG 273
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 263 ATGCGCTTTTGGCAGTGTGATTTGCCAATTACTATTAAGAGGTGAGTCTACTTCCCA 205
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 274 ATATATTGCACTAGATGSCCTTCTGACTTGTGCAATGCAATGATGTAGTACAAATGA 333
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 204 ACCCTTGAATCTGGAGGGGCTTGAATTTGCTTGTCCAAATGAAATGTATGGAAGTGG 145
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 334 CACTGTGCAACTTTGATTTAGGTTTCGAGAGAACTTACACCTTCACACTCAAG 385
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 144 ACCTCTGTGACTTCTGAGACGAGGTGAGAGGCGATGCGAGTTTCCCTC 92
DB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 5
LOCUS      AG046887      655 bp      DNA      linear      GSS 02-NOV-2001
DEFINITION Pan troglodytes DNA, clone: PTB-026022.F, genomic survey sequence.
ACCESSION   AG046887
VERSION     AG046887.1 GI:16583773
KEYWORDS    GSS.
SOURCE      Pan troglodytes (chimpanzee)
ORGANISM    Pan troglodytes
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Euthera; Primates; Catarrhini; Hominidae; Pan.
1
Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
Totoki,Y., Watanabe,H. and Sakaki,Y.
BAC end sequences of Library PTB
2 (bases 1 to 655)
Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
Totoki,Y., Watanabe,H. and Sakaki,Y.
Direct Submission
Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suehiro-chou,Tsukuba-shi, Ibaraki, Japan, 305-8565, Japan;
E-mail:chimpbesgsc.riken.go.jp URL:http://hgp.gsc.riken.go.jp/,
Tel:81-45-503-9111, Fax:81-45-503-9170)
Clones are derived from the chimpanzee BAC library PTB This BAC end
was generated during the R&D process and may have higher chance of
clone tracking errors.
PRIMERS
Sequencing: -21M13
L:BRARY
Vector      : pKS145
R.Site 1    : SacI
R.Site 2    : SacI
Location/Qualifiers
1..655
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="PTB-026022.F"
/sex="male"
/cell_type="lymphoblast"

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[illegible]

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/clone="DKF3p686B1128"  
/tissue_type="human skeletal muscle"  
/dev_stage="adult"
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/lab_host="PH103"
/clone_lib="666" (synonym: h1003)
/notes="vector: pTriplex2; Site_1: SfiI; Site_2: SfiI;
cDNA-collection"
BASE COUNT      174 a      105 c      147 g      162 t
ORIGIN
Query Match      13.3%; Score 66.4; DB 9; Length 585;
Best Local Similarity 58.7%; Pred. No. 0.3;
Matches 160; Conservative 0; Mismatches 116; Indels 3; Gaps 3;
QY 16: TGTGCTAAATTAATTCGCAAGAGTGGCCACACAAATTCCTGCTCATATATATGCGG 220
DB 447 TGAAGTAGATGTTTATAGAGAGGCCACAA-ATTTCCTCTACCTTTCATTCATGTC 389
QY 22: CTTTGGCAATGAGCTTTGCTACTTCTCT-ATCAGATGAGAGCTTATTTCCATATAT 279
DB 388 TTTTGTCTGCTGACTTTGCGGCTTCGCCATCAAGAACTGAGTCCATTCGACCTCGG 329
QY 280 TGCAGTAGATGGCCCTTCGACTGCTTTTGAATGGAATGATAGTACAAATGACACTGT 339
DB 328 TGAATCTGCGCTGGCCCTTCGCTTTCAGCAGTCAATGAAATGGAATGGAATTCAC 269
QY 340 GCAACTTGGATTTAGTTTGGAGAGAACTTACACCTCCACTGACACTCTCTTGGAA 399
DB 268 ACAACTACTAGCTAGTCTCAGTGGCTTCAATTTTCACCTCTCTCTCTCTCTCT 209
QY 400 CCAGATGCAATGTAAGAAGTCAGGGCTATCTCGCTAGAGACATATGT 447
DB 208 AGATCT-TTTGTAAATAGTCTGAGACAGGCTGCTATAGAGATTAGT 162

RESULT 10
LOCUS      A1041146/c
DEFINITION 0969f02.x1 Soares testis_NHT Homo sapiens cDNA clone IMAGE:1542587
3', mRNA sequence.
ACCESSION  A1041146
VERSION     A1041146.1 G:3290340
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
REFERENCE  1. (bases 1 to 477)
AUTHORS   Eukaryota; Vertebrata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
          (bases 1 to 477)
          NC-CGAP http://www.ncbi.nlm.nih.gov/ncgap/.
          National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
          Tumor Gene Index
          Unpublished
          Contact: Robert Strausberg, Ph.D.
          Email: cgapbs@mail.nih.gov
          cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo
          , Ph.D.
          cDNA Library Arrayed by: Greg Lennon, Ph.D.
          DNA Sequencing by: Washington University Genome Sequencing Center
          Clone distribution: NC-CGAP clone distribution information can be
          found through the J.M.A.G.E. Consortium/LLNL at:
          www-bio.lnl.gov/bbrp/image/image.html
          Insert Length: 974      Std Error: 0.00
          Seq primer: -40m13 fwd. ET from Amersham
          High quality sequence stop: 462.
          Location/Qualifiers
            1..477
              /organism="Homo sapiens"
              /mol_type="mRNA"
              /db_xref="taxon:9606"
              /clone="IMAGE:1642587"
              /sex="male"
              /lab_host="PH108"
              /clone_lib="Soares testis_NHT"
              /note="vector: pTriplex2 (Pharmacia) with a modified
              polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
              was prepared from mRNA obtained from Clontech Laboratories
```

```
. Inc., and primed with a Not I - oligo(dT) primer (5'-
TGTACCACTCTAGTGGAGCGCGCCCAATTTTCTTTTCTTTT 3').
Double-stranded cDNA was ligated to Eco RI adapters
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pTri3 vector. Library
went through one round of normalization to cDNA, and was
constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT      126 a      98 c      107 g      146 t
ORIGIN
Query Match      13.1%; Score 65.6; DB 9; Length 477;
Best Local Similarity 62.5%; Pred. No. 0.4;
Matches 132; Conservative 0; Mismatches 74; Indels 2; Gaps 2;
QY 234 CTTTGTCTTCTTCTTCAAGATGTGGAGCTTNTTTCGCATATATTCGACTAGTGG 293
DB 410 CTTTGCACCTCTCCCTTATGAGTGGAGCTTATTCATTCCTTTCGACTGGACTGA 351
QY 294 CTTTGTCTTCTTCTTCAAGATGTGGAGCTTNTTTCGCATATATTCGACTAGTGG 351
DB 350 CTTTGTCTTCTTCTTCAAGATGTGGAGCTTNTTTCGCATATATTCGACTAGTGG 291
QY 352 TTAGGCTTTCGAGAGAACTTACACTTCACCTTCCTTTCGAAACGAGATGCAATG 411
DB 290 TTAGGCTTTCGAGAGAACTTACACTTCACCTTCCTTTCGAAACGAGATGCAATG 231
QY 412 TTAAGAAGTCAAGGCTATCTCTGCTAGAG 439
DB 230 CTAAGGAAGTCAAGGCTATCTCTGCTAGAG 203

RESULT 11
LOCUS      BF29327/c
DEFINITION 152-NT0202 061200-238-c08 NT0202 Homo sapiens cDNA, mRNA sequence.
ACCESSION  BF29327
VERSION     BF29327.1 G:12327455
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
REFERENCE  1. (bases 1 to 474)
AUTHORS   Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
          (bases 1 to 474)
          Dias Neto, E., Garcia Correa, R., Vertovski-Almeida, S., Briones, M.R.,
          Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.P.,
          Goldman, S.H., Carvalho, A.F., Katsukuma, A., Bala, S.S., Simpson, D.H.,
          Brustein, A., de Oliveira, P.S., Bucher, P., Jorgensen, C.V., O'Hare
          , M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and
          Simpson, A.J.
          Shotgun sequencing of the human transcriptome with GRF expressed
          sequence tags
          Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
          20202663
          10737820
          Contact: Simpson A.J.G.
          Laboratory of Cancer Genetics
          Ludwig Institute for Cancer Research
          Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
          Brazil.
          Tel: +55-11-2704922
          Fax: +55-11-2707601
          Email: asimpson@ludwig.org.br
          This sequence was derived from the FAPESP/LICR Human Cancer Genome
          Project. This entry can be seen in the following URL
          http://www.ludwig.org.br/scripts/gethtml2.pl?::=12&t2=12-NT0202-061200-238-c08&t3=2003-12-08&t4=1
          Seq primer: puc 18 forward
          High quality sequence start: 59
          High quality sequence stop: 443.
          Location/Qualifiers
            1..474
              /organism="Homo sapiens"
              /mol_type="mRNA"
```


AUTHORS

Kawai, J., Shiragawa, A., Saibata, K., Yoshino, Y., Itoh, X., Ishii, Y., Arakawa, T., Hara, A., Fukushiro, Y., Komuro, H., Adachi, J., Fukuda, S., Aizawa, K., Izawa, M., Nishi, K., Kiyosawa, H., Kondo, S., Yamanaka, T., Saito, T., Okazaki, Y., Gotohori, T., Bono, H., Kasukawa, T., Saito, R., Kadota, K., Matsuda, H., Ashtukner, V., Batalov, S., Casavant, T., Fleisemann, W., Gaasterland, T., Gissi, C., King, B., Kochiwa, H., Kuehl, P., Lewis, S., Macosio, Y., Nakaide, T., Pesole, G., Rost, D., Quackenbush, J., Schriml, L., Staub, F., Suzuki, R., Temita, M., Wagner, L., Washio, T., Sakai, K., Okido, T., Furuno, M., Aono, H., Baldarelli, R., Barsh, G., Blake, J., Boffelli, D., Bojunga, N., Carinci, P., de Bonaldo, V., Brownstein, M., J., Sait, C., Fletcher, C., Fujita, M., Gariboldi, N., Gustincich, S., Hill, D., Hochmann, M., Hume, D., Kariya, M., Lee, R., H., Lyons, P., Machinomi, S., Mashima, J., Mazzarelli, J., Mochizuki, P., Nordens, P., Ring, B., Ringwald, M., Rodriguez, I., Sakamoto, K., Sakai, H., Sato, K., Schonbach, C., Seya, T., Shibata, Y., Storch, K. F., Suzuki, H., Toyooka, K., Wang, K. H., Weitz, C., Whitaker, C., Williams, L., Wyshaw-Boris, A., Yoshida, K., Hasegawa, Y., Kawai, H., Kohsaki, S., and Hayashizaki, Y.

Functional annotation of a full-length mouse cDNA collection

Nature 409 (5821), 695-699 (2001)

2108566C

11217951

REFERENCE

AUTHORS

The PANTOM Consortium and the RIKEN Genome Exploration Research

Group Phase I & II Team

Analysis of the mouse transcriptome based on functional annotation

of 60,770 full-length cDNAs

Nature 420, 563-573 (2002)

6 (bases 1 to 1335)

JOURNAL

AUTHORS

Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carrinai, P., Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W., Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Hori, F., Imokani, K., Ishii, Y., Itoh, M., Kagawa, T., Kasukawa, T., Katoh, H., Kawai, J., Koyama, Y., Kondo, S., Komuro, H., Kouda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, Y., Nakamura, Y., Nishi, K., Nomura, K., Nuzaki, R., Ogo, Y., Ohsato, N., Okazaki, Y., Saito, R., Saichoh, H., Sakai, C., Sakai, K., Sakazume, N., Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sobabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Takeda, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A., Muramatsu, M., and Hayashizaki, Y.

Direct Submission

Submitted (16-JUL-2001) Yoshinori Hayashizaki, The Institute of

Physical and Chemical Research (RIKEN), Laboratory for Genome

Exploration Research Group, RIKEN Genomic Sciences Center (GSC),

RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Teikoku-Ku, Yokohama,

Kanagawa 230-0045, Japan (E-mail: genome-res@gs.c.riken.go.jp).

URL: <http://genome.gsc.riken.go.jp/>, Tel: 61-45-503-9222.

Fax: 61-45-503-9216

cDNA library was prepared and sequenced in Mouse Genome

Encyclopedia Project of Genome Exploration Research Group in RIKEN

Genomic Sciences Center and Genome Science Laboratory in RIKEN

Division of Experimental Animal Research in Riken contributed to

prepare mouse tissues.

Please visit our web site for further details.

URL: <http://genome.gsc.riken.go.jp/>

URL: <http://fantom.gsc.riken.go.jp/>

FEATURES

source

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CC:42611/C

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

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COMMENT

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 RX116065
 RX116065.1 GI:27890373

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 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo;
 1 (Bases 1 to 497)
 Ebert,D., Reil,O., Hennig,S., Neubert,P., Partsch,R., Peters,M.,
 Radecki,U., Schneider,D. and Korn,B.
 Human UnigeneSet - RZPD3

Unpublished
 Contact: Ina Rolfs
 RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
 77, Neuenheimer Feld 580, D-69120 Heidelberg, Germany
 RZPD; IMAGP998E035152
 RZPDHIS; I.M.A.G.E. cDNA clone collection
 Human UnigeneSet - RZPD3 (RZPDHIS No.342)
 http://www.rzpd.de/CloneCards/cgi-bin/showlib.pl.cgi?response=libNo:372 Contact: Ina Rolfs
 RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
 Neuenheimer Weg 5, D-14059 Berlin, Germany
 Tel: +49 30 32639 101
 Fax: +49 30 32639 111
 www.rzpd.de

This clone is available royalty-free from RZPD;
 contact RZPD (clone@rzpd.de) for further information. Seq primer:
 M33r, Primer sequence: TTCACAGGAACAGCTATGAC.

FEATURES

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 vitro. Following RAP purification, this DNA was used as
 tracer in a subtractive hybridization reaction. The driver
 was PCR-amplified cDNAs from pools of 5,000 clones made
 from the same 3 libraries. The pools consisted of
 I.M.A.G.E. clones 297480-32087, 682632-687239,
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 Soares and M. Fatima Borralho."
 BASE COUNT 154 a 98 c 120 g 125 t

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Patent: WO 01/19198-A 1 15-MAR-2003;
CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (CNRS) (FR)
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RS246003

LOCUS RS246003 110000 bp DNA linear PRI 08-MAR-2000

DEFINITION Homo sapiens Spast gene for spastin protein.

ACCESSION U246003

VERSION A246003.1 GI:6273492

KEYWORDS Spast gene; spastin protein; SPG4-linked hereditary spastic paraplegia.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1

AUTHORS Hazan,J., Ponkrecht,N., Mavel,D., Paternotte,C., Samson,D., Attiguenave,F., Davoine,C., Cruaud,C., Durr,A., Wincker,P., Brettier,P., Cattolico,L., Barbe,V., Burgunder,J.M., Parfrey,N., Brice,A., Fric,A., Fontaine,B., Prud'homme,J., Weissbach,J., Durr,A. and Hazan,J.

TITLE Spastin, a new AAA protein, is altered in the most frequent form of autosomal dominant spastic paraplegia

JOURNAL Nat. Genet. 23 (3), 296-303 (1999)

MEDLINE 20055425

PUBMED 10610178

REFERENCE 2

AUTHORS Ponkrecht,N., Mavel,D., Byrne,P., Davoine,C., Cruaud,C., Beetsch,D., Samson,D., Coutinho,P., Hutchinson,V., McMonagle,P., Burgunder,J., Fartagione,A., Heinzel,O., Felix,J., Deufel,T., Parfrey,N., Brice,A., Fontaine,B., Prud'homme,J., Weissbach,J., Durr,A. and Hazan,J.

TITLE Spectrum of SPG4 mutations in autosomal dominant spastic paraplegia

JOURNAL Hum. Mol. Genet. 9 (4), 637-644 (2000)

MEDLINE 20164302

PUBMED 10693187

REFERENCE 3

AUTHORS Genoscope.

TITLE Direct Submission

JOURNAL Submitted (17-JUN-1999) Genoscope, Genoscope - Centre National de Sequencage, BP 191, EVRY 91006, FRANCE

COMMENT E-mail : seqref@genoscope.cns.fr Web : www.genoscope.cns.fr. The sequence is the result of the assembly of 2 BAC clones: R-336p14 and 563N4, respectively from RPC1-11 and CTRB_97A_SKB library.

FEATURES

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Matches 501; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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VERSION AL121658.4 GI:20218783
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SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
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AUTHORS Hazan, C., Fonknechten, N., Mavel, D., Paternotte, C., Sanson, D.,
Artiguenave, F., Davoine, C.S., Cruaud, C., Dur, A., Winkler, P.,
Brotier, P., Catolico, B., Barbe, V., Burgunder, J.M.,
Prud'Homme, J.F., Brice, A., Fontaine, B., Heilig, R. and
Weissenbach, J.
TITLE Spastin, a novel AAA protein, is altered in the most frequent form
of autosomal dominant spastic paraplegia
JOURNAL Nat. Genet. (1999) in press
REFERENCE 2 (bases 1 to 142692)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (18-APR-2002) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- web : www.genoscope.cns.fr)
COMMENT On Apr 19, 2002 this sequence version replaced gi:202160242.
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Query Match 100.0% Score 501; DB 9; Length 142692;
Best Local Similarity 100.0% Pred No. 2,7e-116;
Matches 501; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTATTGCTGAGTTCCTTAATCTGAGTCTTGAATCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 60
DB 31259 GGTATTGCTGAGTTCCTTAATCTGAGTCTTGAATCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 31258
QY 61 AGTTGCTGAGTTCCTTAATCTGAGTCTTGAATCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 120
DB 31319 AGTTGCTGAGTTCCTTAATCTGAGTCTTGAATCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 31318
QY 121 GGTCAATTTTGAATAGGTGTGCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 160
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DB 31379 GGTCAATTTTGAATAGGTGTGCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 31378
QY 181 GGTATTGAGTTCCTGAGAGTCTTGAATCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 240
DB 31439 GGTATTGAGTTCCTGAGAGTCTTGAATCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 31438
QY 241 GGTATTGAGTTCCTGAGAGTCTTGAATCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 300
DB 31499 GGTATTGAGTTCCTGAGAGTCTTGAATCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 31498
QY 301 TTGTGTGGAGTCTTAAGTCTTCTTGAATCTGCTTGAAGTCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 360
DB 31559 TTGTGTGGAGTCTTAAGTCTTCTTGAATCTGCTTGAAGTCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 31558
QY 361 CTCCTGTATTTGGTGCATATATATTAGAGTCTGCTTGAAGTCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 420
DB 31619 CTCCTGTATTTGGTGCATATATATTAGAGTCTGCTTGAAGTCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 31618
QY 421 TTAGAGTATATAGTGGGCTCTTCTTGAATCTGCTTGAAGTCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 480
DB 31679 TTAGAGTATATAGTGGGCTCTTCTTGAATCTGCTTGAAGTCTGAGAGTGGGGTGGTGAAGTCTCCAGTATTA 31678
QY 481 TATCAGAGAGTGGATTGCAA 501
DB 31739 TATCAGAGAGTGGATTGCAA 31738

RESULT 4
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LOCUS Homo sapiens chromosome 2 clone RP11-78E13, WORKING DRAFT SEQUENCE,
DEFINITION 7 unordered pieces.
ACCESSION AC011232
VERSION AC011232.7 GI:13270720
KEYWORDS HTS; HTGS, PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 185281)
AUTHORS Waterston, R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 185281)
AUTHORS Waterston, R.H.
TITLE Direct Submission
JOURNAL Submitted (04-OCT-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
COMMENT On Mar 10, 2001 this sequence version replaced gi:9795911.
ORIGIN
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H NH0378E13
Center summary statistics
Sequencing vector: M13, 57%
Chemistry: Dye-terminator Big Dye; 50% of reads
Assembly program: Phrap; version 0.990119
Consensus quality: 182798 bases at least Q40
Consensus quality: 183538 bases at least Q30
Consensus quality: 184045 bases at least Q20
Insert size: 18000; agarose-gel
Insert size: 184681; sum-of-contigs
Quality coverage: 7.07 in Q20 bases; agarose-gel
Quality coverage: 7.33 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently
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REFERENCE 3 (bases 1 to 173106)
 AUTHORS Worley,K.C.
 TITLE Direct Submission
 JOURNAL Submitted (31-OCT-2003) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 REFERENCE 4 (bases 1 to 173106)
 AUTHORS Worley,K.C.
 TITLE Direct Submission
 JOURNAL Submitted (01-DEC-2001) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 COMMENT On Oct 31, 2003: this sequence version replaced g1:26418054.
 INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email
gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
 entire insert of this clone. Overlapping regions of clones are only
 sequenced and submitted once, so the sequence for the remainder of
 the insert may be found in the record for the adjacent clones.
 Overlapping clones are noted at the beginning and end of the
 features listing.

ANNOTATION OF FEATURES:
 STSs are identified using EPCR (Genome Res. 7:541-550) searches
 of a local database that includes entries from dbSTS, GDB, and
 local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green,
 unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST
 (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-14) to the
 EST and cDNA sequences. Genes demonstrate at least two exons
 flanked by consensus splice sites that maintained sequence
 continuity across the splice junctions. Sequences that are not
 identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum
 standard of double strand coverage with a minimum of 2 clones and 2
 reads with no ambiguities or 2 chemistries with a minimum of 2
 clones and 3 reads with no ambiguities. If the sequence quality for
 a region does not meet this standard, it will be indicated in the
 annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality
 standards. Estimated error rate less than 1 per 10,000 bases.
 Reports of lowest quality individual bases and measures of base
 quality are listed below. Description of the metrics can be found
 at UR2.

http://gc.bcm.tmc.edu:8088/quality/info/genbank_annotation.html

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 /function="clone overlap"
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 2220..2342
 /rpt_family="MIR"
 2951..3613
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 4275..4612
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Query Match 88.9%; Score 445.6; DB 9; Length 173106;
 Best Local Similarity 93.4%; Pred. NO. 4.3e-104;
 Matches 482; Conservative 0; Mismatches 19; Indels 15; Gaps 1;

Collins, S., Collymore, A., Cooke, P., Dearellano, K., Dewar, K.,
 Diaz, S., Dodge, S., Faro, S., Ferreira, P., FitzHugh, K., Gage, D.,
 Gagan, J., Gardyna, S., Ginde, S., Goyette, M., Graham, J.,
 Grand-Pierre, K., Hagos, B., Heaford, A., Horton, L., Hulme, W.,
 Ilev, I., Johnson, R., Jones, C., Karas, A., Lapocque, K.,
 Lamazares, R., Lardes, T., Lebeck, C., Levine, R., Liu, S.,
 MacLean, C., MacDonald, P., Marquis, N., Matthews, C., McCarthy, M.,
 McKernan, K., McPheters, R., Veldman, J., Menendez, J.,
 Minerva, V., Murphy, J., Naylor, S., Nguyen, C., North, C.,
 Norman, C.H., O'Connor, T., O'Donnell, P., O'Malley, D., Oliver, J.,
 Petersen, K., Phukhang, P., Pierron, K., Pollara, V., Raymond, C.,
 Retta, R., Rietack, X., Riley, R., Rise, C., Rogov, P., Roman, J.,
 Rosetti, M., Roy, A., Santos, A., Schauer, S., Schupbach, R., Seaman, S.,
 Severy, P., Sougniez, C., Spencer, B., Stange-Thorann, K.,
 Stojanovic, N., Strauss, N., Subramanian, A., Talamas, C., Tesfaye, S.,
 Theodore, C., Travets, X., Travis, N., Tricollon, J., Vassiliev, H.,
 Viel, R., Vo, A., Wilson, B., Wolz, W., Wyman, D., Ye, W.J., Young, G.,
 Zalcoun, C., Zembek, S., Zimmer, A. and Zody, X.

Direct Submission
 Submitted (02-MAY-2001) Whitehead Institute/MIT Center for Genome
 Research, 325 Charles Street, Cambridge, MA 02141, USA
 On May 1, 2001 this sequence version replaced G12313933.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center -----
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence.submissions@genome.wi.mit.edu
 ----- Project Information -----
 Center Project name: L2082
 Center Clone name: 219_J_21

FEATURES

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Lamazares, P., Landers, T., Lenockzy, J., Levine, R., Liu, G., Maclean, C., Macdonald, P., Major, J., Margolis, M., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., McPherson, P., Vaidyan, J., Veneus, L., Xhova, T., Mienga, V., Murphy, T., Nay, G., Nguyen, C., Norbu, C., Norman, C.H., O'Connor, T., O'Connell, P., O'Neill, C., Oliver, J., Peterson, K., Prunkhang, P., Pierre, N., Polara, V., Raymond, C., Retta, R., Riback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schaefer, S., Schupbach, P., Sevan, K., Severy, P., Spencer, B., Stange, R., Stange, R., Stange, R., Strauss, K., Subramanian, A., Talamas, J., Testa, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigg, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W., Young, G., Zalcov, J., Zemlek, L., Zimmer, A., and Zody, N.

Submitted (01-DEC-2001): Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Nov 29, 2001 this sequence version replaced g11604527.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RN/RepeatMasker.html>

Center: Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WPR

Web site: <http://www.seq.wi.mit.edu>

Contact: sequence_submissions@genome.wi.mit.edu

Project Information

Center project name: 57841

Center C-line name: 227_F_6

Location/Qualifiers

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Query Match 89.3% Score 442.4; DB 9; Length 161644;
Best Local Similarity 93.0%; Pred. No. 2.9e-103;
Matches 480; Conservative 0; Mismatches 21; Indels 15; Gaps 1;

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b 142139 AGTTTGTGTAATTCCTGTTCTTTTACATTTGCTGAGGAGTGTGTTAGTTCACCACTATGT 142196

Y 121 GGTCAATTTTGGATAGTGTGGTGTGGTCTGAGAGAGATGTATATCTTGTGATTTGG 180
b 142199 GGTCAATTTTGGATAGTGTGGTGTGGTCTGAGAGAGATGTATATCTTGTGATTTGG 42228

Y 161 GGTTCAGAGTCTCTAGATGTCATTAGGTCACCTGCTGAGAGTGTGAGTTCAGTTCCT 240
b 142259 GGTTCAGAGTCTCTAGATGTCATTAGGTCACCTGCTGAGAGTGTGAGTTCAGTTCCT 42318

Y 241 GGA-----TCTGCTTTTCTGATCTGCTTAAATATGACAGTGTGCTTGA 285
b 142319 GGAATATCTTTTAACTTCTGCTTAAATATGACAGTGTGCTTGA 42378

Y 286 AGTCTCCAGCATATTGCTGTGAGTCTAGTCTCTTCTAGATCTCTTACAGAGTCTGT 345
b 142379 AGTCTCCAGCATATTGCTGTGAGTCTAGTCTCTTCTAGATCTCTTACAGAGTCTGT 42438

Y 346 TTATGAATCGGGTGCCTCTGATTTGGTGCATATATATTAGATAGTTCCTCTCT 405
b 142439 TTATGAATCGGGTGCCTCTGATTTGGTGCATATATATTAGATAGTTCCTCTCT 42498

Y 406 GTTGAATGATCCCTTAGCATATATGATGGCTCTTCTGCTCTCTTCTGATTTCTG 465
b 142499 GTTGAATGATCCCTTAGCATATATGATGGCTCTTCTGCTCTCTTCTGATTTCTG 42559

Y 466 GTTAAAGTCTCTTTATCAGAGAGTTCGATTGCA 501
b 142559 GTTAAAGTCTCTTTATCAGAGAGTTCGATTGCA 42594

RESULT 12
NS01DVT 169505 bp DNA linear PRI 01-JUN-2003
CCUS Human chromosome 14 DNA sequence BAC R-149123 of library RPCL-11
EFINITION from chromosome 14 of Homo sapiens (human), complete sequence.
CCSS:CN AL132987.4 GI:14284936
ERS:CN HTG: HTGS ACTIVEFIN
EYMCRDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 169505)
Heilig, R., Petit, J.L., Vico, V., Dasilva, C., Robert, C., Wierker, P.,
Brottier, P., Catalicio, L., Barbe, V., Pelletier, E., Artiguerave, P.,
Jevy, M., Eckenberg, R., Bruls, T., Debernardinis, V., Crnaud, C.,
Gyapay, G., Saurin, W. and Weissbach, J.
Sequencing of the human chromosome 14
Unpublished
2 (bases 1 to 169505)
Genoscope.
AUTHORS Direct Submission
TITLE Submitted (31-MAY-2001) Genoscope - Centre National de Sequencage :
JOURNAL BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
```

On Jun 3, 2001 this sequence version replaced g1:8217897.
----- Genoscope
Center: Genoscope / Centre National de Sequencage
Web site: http://www.genoscope.cns.fr/
Contact: SeqRef@genoscope.cns.fr

The following BAC sequence is oriented from the T7 to the SP6 end.
Upstream BAC (overlapping the T7 end) : R-862015 (AC=Alu17192)
Downstream BAC (overlapping the SP6 end) : R-807016 -----

Summary Statistics
Assembly program: Phrap; version 2.0
Quality coverage: 8.35x in Q20 bases; sum-of-contigs -----

Overall quality chart :
Range : bases
C 1 - 9 :
10 - 19 : 3
20 - 29 : 6
30 - 39 : 43
40 - 49 : 716
50 - 59 : 1710
60 - 69 : 5293
70 - 79 : 19513
80 - 89 : 59301
90 - 99 : 82920

Percentage of bases with a quality value >= 40 : 99 %

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/chromosome="14"
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/note="matching EMSL:F01540
RHdb:RH26869
dbSTS:STS6768
Identified using the e-PCR software (G. Schuler)"
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dbSTS:STS47040
Identified using the e-PCR software (G. Schuler)"

BASE COUNT 47640 a 36003 c 37377 g 48485 t
ORIGIN

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Best Local Similarity 93.0%; Pred. No. 2.9e-103;
Matches 480; Conservative 0; Mismatches 21; Indels 15; Gaps 1;

QY 1 GGTITTCGTCAGGTTCTTAACTCTGAGTCTTACTGATTTGATTCGAGAGAC 60
Db 29793 GGTITTCGTCAGGTTCTTAACTCTGAGTCTTACTGATTTGATTCGAGAGAC 29852

QY 61 AGTTTGTGTAATTCCTGTTCTTTTACATTTGCTGAGGAGTGTGTTAGTTCACCACTATGT 120
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QY 121 GGTCAATTTTGGATAGTGTGGTGTGGTCTGAGAGAGATGTATATCTTGTGATTTGG 180
Db 29913 GGTCAATTTTGGATAGTGTGGTGTGGTCTGAGAGAGATGTATATCTTGTGATTTGG 29972

QY 161 GGTTCAGAGTCTCTAGATGTCATTAGGTCACCTGCTGAGAGTGTGAGTTCAGTTCCT 240
Db 29973 GGTTCAGAGTCTCTAGATGTCATTAGGTCACCTGCTGAGAGTGTGAGTTCAGTTCCT 30032

QY 241 GGA-----TCTGCTTTTCTGATCTGCTTAAATATGACAGTGTGCTTGA 285
Db 30032 GGAATATCTTTTAACTTCTGCTTAAATATGACAGTGTGCTTGA 30090

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y 346 TTATGAATCTGGTGGTCCCTGCTATGTTGGTGCATATATATTAGATAGTATAGTCTCTCT 405
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RESULT 13
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DEFINITION Homo sapiens 12 BAC RP11-714X16 (Rosenfeld Park Cancer Institute Human BAC Library) complete sequence.
ACCESSION AC080012
VERSION AC080012.20 GI:14290372
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
1 (bases 1 to 99966)
Muzny, D.M., Adams, C., Adio-Oduola, B., Ali-Osman, F.R., Allen, C.,
Ashbrook, S.B., Amaral, H.C., Aze, R.R., Banks, J.T., Barbara, J.,
Benton, J., Bimaga, K., Blackenburg, K., Bonnin, D., Bouck, J.,
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Carter, M., Cavazos, S.R., Chacko, J., Chavez, D., Chen, G., Chen, R.,
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Cox, C., Coyne, M.D., Dabney, S.R., David, R., Davila, M.D., Davis, C.,
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Dugan-Rocha, S., Durbin, K.J., Earhart, C., Edgar, D., Edwards, C.C.,
Elhaj, C., Eerling, S., Escotto, M., Falis, T., Ferraguto, D.,
Flagg, N., Ford, J., Foster, P., Frantz, P., Gabris, A., Gao, J.,
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Hodgson, A., Hogues, M., Holloway, C., Hollins, B., Horst, P.,
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Zorrilla, S., Zucherlapati, R. and Gibbs, R.

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Direct Submission
Unpublished
2 (bases 1 to 99966)
Worley K.C.
Direct Submission
Submitted (23-SEP-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 99966)
Worley K.C.
Direct Submission
Submitted (05-JUN-2001) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 99966)
Worley K.C.
Direct Submission
Submitted (07-MAY-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
5 (bases 1 to 99966)
Worley K.C.
Direct Submission
Submitted (31-MAY-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Jun 5, 2001 this sequence version replaced gi:4017474.
INFORMATION: http://www.hgsc.bcm.tmc.edu/ or email
gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
entire insert of this clone. Overlapping regions of clones are only
sequenced and submitted once, so the sequence for the remainder of
the insert may be found in the record for the adjacent clones.
Overlapping clones are noted at the beginning and end of the
Features listing.

ANNOTATION OF FEATURES:
STSS are identified using ePCR (Genome Res. 7:541-550) searches
of a local database that includes entries from dbSTS, GDB, and
local mapping efforts.
Repeats are identified using RepeatMasker (A. Smit and P. Green,
unpublished.) for Human and Mouse sequences.
Genes and Region of sequence similarity are identified by BLAST
(Nuc. Acids Res. 25:3389-3402) similarity expect < 1e-34 to the
EST and cDNA sequences. Genes demonstrate at least two exons
flanked by consensus splice sites that maintained sequence
continuity across the splice junctions. Sequences that are not
identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum
standard of double strand coverage with a minimum of 2 clones and 2
reads with no ambiguities or 2 chemistries with a minimum of 2
clones and 3 reads with no ambiguities. If the sequence quality for
a region does not meet this standard, it will be indicated in the
annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality
standards - estimated error rate less than 1 per 10,000 bases.
Reports of lowest quality individual bases and measures of base
quality are listed below. Description of the metrics can be found
at URL:
http://gc.bcm.tmc.edu:2086/quality.info/genbank.annotation.html.

QUALSTAT-REPORT-----
----- Summary Statistics -----
Contig length: 73346
Phrap values in estimate: 73205
Average error rate (BCM-Phrap estimate): 8.64574e-05
Fraction of Phrap values less than 40: 0.00487652
Number of consensus changing edits: 0

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ambiguous, there is an annotation using the 'unsure' feature key. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: EMBL, EMBL; SW, SwissProt; Tr, TrEMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep. This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/ChrX>. RPL1-45J1 is from the library RPL1-1.1 constructed by the group of Peter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>. VECTOR: pBACel.5

IMPORTANT: This sequence is not the entire insert of clone RPL1-45J1. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.

The true left end of clone RP6-5234 is at 91766 in this sequence. The true right end of clone RPL1-525J14 is at 100 in this sequence.

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Best Local Similarity 92.8%; Pred. No. 7; Set 103;
Matches 479; Conservative 0; Mismatches 22; Indels 15; Gaps 1;
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2y 121 GTCATTTTGGAAATAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
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2b 77178 AGCTTCCAGATATTATTGGTGGAGTCTTAAGTCTCTTGGAGTGGTGGTGGTGGTGG 77119
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2b 77116 TTATGAATCTGGTCTCTGATTTGGGTGCATATATATTTAGGATATAGTCTTCTT 77053
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2b

RESULT 15

AY306744
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ACCESSION
VERSION
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REFERENCE
AUTHORS
TITLE
JOURNAL
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Job time : 1390.29 secs

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481	TATCAGACAGATTGGATTGCCAA	501
50480	TATCAGACAGATTGGATTGCCAA	50500
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X	antiallergic; hepatotropic; antidiabetic; antiinflammatory; antitumor;	
X	vulvarly; anticonvulsant; antibacterial; antifungal; antiparasitic;	
X	cardiac; gene therapy; cancer; immune disorder; cardiovascular disorder;	
X	neurological disease; infection; human; secreted protein; ss.	
S	Homo sapiens.	
N	W0230055371-A1.	
D	21-SEP-2000.	
X	16-MAR-2000; 2000WC-US06783.	
X	18-MAR-1999; 990S-0125059.	
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RESULT: 3
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Human; secreted protein; diagnosis; sclerosis; immunomodulatory; antisclerotic;
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immunostimulant; cytostatic; cardiac; vascular; anti-angiogenic;
ophthalmological; neuroprotectant; neurotropic; anticonvulsant; vulneary;
antialzheimers; antiparkinsonian; antimicrobial; immune disorder;
multiple sclerosis; systemic lupus erythematosus; HIV infection;
hyperproliferative disorder; cancer; Gaucher's disease; wound healing;
cardiovascular disease; Schmitz syndrome; Chaga's cardiomyopathy;
corneal arteriosclerosis; angiogenic disorder; diabetic retinopathy;
corneal graft neovascularisation; neurological disorder; regeneration;
Huntington's chorea; Alzheimer's disease; Parkinson's disease;
infectious disease; chemotaxis; ss.

XX	Homo sapiens.
OS	WC20007653C-A.
XX	21-DEC-2000.
PV	
XX	
PD	

```
XX CC 11-JUN-2003; 2000MC-US14933.
XX PF
XX CC 11-JUN-1999; 99US-2138572.
XX CC
XX CC (HUMA-3) HUMAN GENOME SCI INC.
XX PA (ROSE/J) ROSEN C A.
XX PT
XX PI Rosen CA, Ruben SM, Komatsoulis GA;
XX CC
XX PF
XX CC WFI: 2001-0711-47/2a.
XX CC P-PDB; A864888.
XX CC
XX CC Nucleic acids encoding 42 human secreted polypeptides, useful for
XX PT preventing, diagnosing and/or treating e.g. cancers, Parkinson's
XX PT disease and diabetic retinopathy.
XX CC
XX CC Claim 1: Page 453-455; 554pp; Eng:1sh.
XX CC
XX CC The polynucleotide sequences given in AAF3321.3 to AAF3325.1 encode the
XX CC human secreted proteins given in AAB64882 to AAB64930. AAB64931 to
XX CC AAB64991 represent human secreted polypeptide sequences and proteins
XX CC homologous to them, which are given in the exemplification of the present
XX CC invention. Human secreted proteins have activities based on the tissues
XX CC and cells the genes are expressed in. Examples of activities include:
XX CC immunomodulatory; antisclerotic; dermatological; immunosuppressive;
XX CC antiinflammatory; anti-HIV; immunostimulant; cytostatic; cardiac;
XX CC vascular; antimicrobial; anti-angiogenic; ophthalmological;
XX CC neuroprotectant; anticonvulsant; nootropic; antialzheimers;
XX CC antiparkinsonian; and vulnery. The polynucleotides and polypeptides can
XX CC be used in the prevention, diagnosis and treatment of diseases associated
XX CC with inappropriate polypeptide expression. Disorders that may be
XX CC prevented, diagnosed and/or treated by the above methods include immune
XX CC disorders (e.g. multiple sclerosis, systemic lupus erythematosus and
XX CC human immuno-deficiency virus (HIV) infections), hyperproliferative
XX CC disorders (e.g. cancers and Gaucher's disease), cardiovascular diseases
XX CC (e.g. Scimitar syndrome, Chaga's cardiomyopathy and coronary
XX CC arteriosclerosis), angiogenic disorders (e.g. corneal graft
XX CC neovascularisation and diabetic retinopathy), neurological disorders
XX CC (e.g. Huntington's chorea, Alzheimer's disease and Parkinson's disease),
XX CC infectious diseases and/or for promoting wound healing, regeneration and
XX CC /or chemotaxis. AAF33204 to AAF33212 and AAB64881 represent sequences
XX CC used in the exemplification of the present invention.
XX CC
XX CC Sequence 5065 BP; 980 A; 948 C; 1046 G; 2017 T; 74 other;
XX CC
XX CC Query Match 87.0%; Score 436; DB 22; Length 5065;
XX CC Best Local Similarity 91.7%; Pred. NC. 4.4e-108;
XX CC Matches 473; Conservative 4; Mismatches 24; Indels 15; Gaps 1;
XX CC
XX CC 1 GGTGTTGGTGGAGTTCTTAATCTGAGTCTAGTCTAGTCTAGTCTGGCTGAGAC 60
XX CC 3355 GGTGTTGGAGTGGAGTTCTTAATCTGAGTCTAGTCTAGTCTAGTCTGGCTGAGAG 3454
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XX CC 61 AGTTGTTGTAATTTCTGTTCTTCTACATTTGCTGAGGAGTCTTGTAGTCCAACTATCT 120
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XX CC
XX CC 121 GGTCAATTTGGATAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
XX CC 3475 GGTCAATTTGGATAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 3534
XX CC
XX CC 181 GGTGTTAGAGTTCTGTAGATGTCGTATAGTTCAGTCTGGTGGAGAGTGGTGGTGGT 240
XX CC 3535 GGTGTTAGAGTTCTGTAGATGTCGTATAGTTCAGTCTGGTGGAGAGTGGTGGTGGT 3594
XX CC
XX CC 241 GG-----ATCTGCTCTTGTGATCTGTCTAATATTGACAGTGGGAGTTCTGA 265
XX CC 3595 GGTATATCTGTTTACTTCTGCTGCTGATCTGTCTAATATTGACAGTGGGAGTTCTGA 3654
XX CC
XX CC 286 AGTCTCCAGTATTATTGTGTGGAGTCTAAGTCTCTTTGTAGTCTCTAGGAGTTGCT 345
XX CC 3655 AGTCTCCAGTATTATTGTGTGGAGTCTAAGTCTCTTTGTAGTCTCTAGGAGTTGCT 3714
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PT variation: /standard_name= "Single nucleotide polymorphism"
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Query Match: 97.0%; Score 436; DB 24; Length 1691080;
 Best Local Similarity 92.2%; Pred. No. 2.5e-107;

Matches 476; Conservative 0; Mismatches 25; Indels 15; Gaps 1;	
QY	1 GATTTGGTGAATTCCTTAATTCCTGAGTTCAGTTTCACTGCTGGCTGAGAGAC 60
Db	12249 GATTTGGTGAATTCCTTAATTCCTGAGTTCAGTTTCACTGCTGGCTGAGAGAC 12308
QY	61 AGTTTGTGTAATTCCTGTTTCTTACATTTCTGAGAGTSCCTTTAGTCCAACTAGT 120
Db	12309 AGTTTGTGTAATTCCTGTTTCTTACATTTCTGAGAGTSCCTTTAGTCCAACTAGT 12368
QY	121 GTCATATTTGGAATAGGTGCTGTTGCTGCTGAGAGAGATATATATCTTGTGATTGG 180
Db	12369 GTCATATTTGGAATAGGTGCTGTTGCTGAGAGAGATATATATCTTGTGATTGG 12428
QY	181 GATTAGAGTCTGTAGATGCTCAATAGTCCACTGCTGAGAGAGTCTGAGTTTCT 240
Db	12429 GATTAGAGTCTGTAGATGCTCAATAGTCCACTGCTGAGAGAGTCTGAGTTTCT 12488
QY	241 GGA-----TCTGCTGTTGATCTGTCTAATATGACAGTGGGGTGTGA 285
Db	12489 GGAATCCTGTTAATCTTCTGATGGATCTGCTAATGTTGACAGTGGGGTGTGA 12548
QY	286 AGTCTCCAGTATTATGCTGGAGTCAATCTCTTTGTAGTCTTTAGGAGTTTCT 345
Db	12549 AGTCTCCAGTATTATGCTGGAGTCAATCTCTTTGTAGTCTTTAGGAGTTTCT 12608
QY	346 TTATGAATCTGGTCTCTGTAATTTGGTGGATATATATTTAGGATAGTCTTTCT 405
Db	12609 TTATGAATCTGGTCTCTGTAATTTGGTGGATATATATTTAGGATAGTCTTTCT 12668
QY	406 GTTGAATTGATCCCTTTAGCATATATGATGCTCTCTTGTCTTTGATCTTTGTG 465
Db	12669 GTTGAATTGATCCCTTTATCATATGTAATGCTCTCTTGTCTTTGATCTTTGTG 12728
QY	466 GTTTAAAGTCTTTTATCAGAGAGTTGATTCGAA 501
Db	12729 GTTTAAAGTCTTTTATCAGAGAGTTGATTCGAA 12764
RESULT 5	
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IC	AA135853 standard; DNA: 9192 BP.
XX	
AC	AA135853;
XX	
DT	08-JAN-2002 (first entry:
XX	
DE	Human musculoskeletal system related polynucleotide SEQ ID NO 2116.
XX	
KW	Cytostatic; immunosuppressive; neurotropic; neuroprotective; antiviral;
KW	antiallergic; hepatotropic; antidiabetic; anti-inflammatory; antitumor;
KW	vulnerable; anticonvulsant; antibacterial; antifungal; antiparasitic;
KW	cardiac; gene therapy; cancer; immune disorder; cardiovascular disorder;
KW	neurological disease; infection; human; secreted protein;
KW	-musculoskeletal system; ds.
XX	
XX	Homo sapiens.
XX	
XX	WC200155367-A1.
XX	
XX	02-AUG-2001.
XX	
XX	17-JAN-2001; 2001WO-US01338.
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XX	31-JAN-2000; 2000US-0179065.
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PR	24-FEB-2000; 2000US-0184564.
PR	02-MAR-2000; 2000US-0186350.
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PR	17-MAR-2000; 2000US-0190076.
PR	18-MAR-2000; 2000US-0198123.
PR	19-MAY-2000; 2000US-0205515.
PR	07-JUN-2000; 2000US-0209467.

PR	26-JUN-2000; 2000US-0214886.
PR	30-JUN-2000; 2000US-0215335.
PR	07-JUL-2000; 2000US-0216647.
PR	11-JUL-2000; 2000US-0216880.
PR	11-JUL-2000; 2000US-0217487.
PR	11-JUL-2000; 2000US-0217496.
PR	14-JUL-2000; 2000US-0218290.
PR	26-JUL-2000; 2000US-0220963.
PR	26-JUL-2000; 2000US-0220964.
PR	14-AUG-2000; 2000US-0224518.
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PR	14-AUG-2000; 2000US-0225268.
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PR	14-AUG-2000; 2000US-0225447.
PR	14-AUG-2000; 2000US-0225757.
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PR	18-AUG-2000; 2000US-0226279.
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PR	22-AUG-2000; 2000US-0227182.
PR	23-AUG-2000; 2000US-0227509.
PR	30-AUG-2000; 2000US-0228524.
PR	01-SEP-2000; 2000US-0229287.
PR	01-SEP-2000; 2000US-0229343.
PR	01-SEP-2000; 2000US-0229344.
PR	01-SEP-2000; 2000US-0229345.
PR	01-SEP-2000; 2000US-0229509.
PR	01-SEP-2000; 2000US-0229513.
PR	06-SEP-2000; 2000US-0230437.
PR	06-SEP-2000; 2000US-0230438.
PR	06-SEP-2000; 2000US-0231242.
PR	06-SEP-2000; 2000US-0231243.
PR	08-SEP-2000; 2000US-0231344.
PR	08-SEP-2000; 2000US-0231413.
PR	08-SEP-2000; 2000US-0231414.
PR	08-SEP-2000; 2000US-0232080.
PR	08-SEP-2000; 2000US-0232081.
PR	12-SEP-2000; 2000US-0231968.
PR	14-SEP-2000; 2000US-0232397.
PR	14-SEP-2000; 2000US-0232398.
PR	14-SEP-2000; 2000US-0232399.
PR	14-SEP-2000; 2000US-0232400.
PR	14-SEP-2000; 2000US-0232401.
PR	14-SEP-2000; 2000US-0233063.
PR	14-SEP-2000; 2000US-0233064.
PR	14-SEP-2000; 2000US-0233065.
PR	21-SEP-2000; 2000US-0234223.
PR	21-SEP-2000; 2000US-0234274.
PR	25-SEP-2000; 2000US-0234997.
PR	25-SEP-2000; 2000US-0234998.
PR	26-SEP-2000; 2000US-0235484.
PR	27-SEP-2000; 2000US-0235834.
PR	27-SEP-2000; 2000US-0235936.
PR	29-SEP-2000; 2000US-0236127.
PR	29-SEP-2000; 2000US-0236167.
PR	29-SEP-2000; 2000US-0236368.
PR	29-SEP-2000; 2000US-0236369.
PR	29-SEP-2000; 2000US-0236370.
PR	02-OCT-2000; 2000US-0236802.
PR	02-OCT-2000; 2000US-0237037.
PR	02-OCT-2000; 2000US-0237038.
PR	02-OCT-2000; 2000US-0237039.
PR	02-OCT-2000; 2000US-0237040.
PR	13-OCT-2000; 2000US-0239935.
PR	13-OCT-2000; 2000US-0239937.
PR	20-OCT-2000; 2000US-0240960.
PR	20-OCT-2000; 2000US-0241221.
PR	20-OCT-2000; 2000US-0241785.


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08-DEC-2000; 2000US-251836P.
08-DEC-2000; 2000US-251836P.
09-DEC-2000; 2000US-251836P.
{ROSE/; ROSEN C A.
{RUSE/; RUBEN S M.
{BARA/; BARASH S C.
Rosen CA, Ruben SM, Barash SC;
WPI; 2003-126199/12.
Isolated nucleic acid molecules encoding musculoskeletal system
associated polypeptides, useful for detecting disorders, e.g. cancer -
Disclosure; SEQ ID NO 2219; 321pp; English.
The invention describes an isolated nucleic acid molecule comprising a
sequence encoding musculoskeletal system associated polypeptides useful
for detecting disorders, e.g., cancer or cancer metastases, in animals
or humans. The nucleic acid stimulates re-vascularisation of ischaemic
tissues associated with conditions such as thrombosis, arteriosclerosis,
and other cardiovascular conditions; treats wounds due to injuries,
burns, post-operative tissue repair, and ulcers; stimulates angiogenesis
and limb regeneration; stimulates neuronal growth; can treat and prevent
neuronal damage occurring in certain disorders or neurodegenerative
conditions, such as, Alzheimer's disease, Parkinson's disease, and
AIDS-related complex; stimulates chondrocyte growth, thus they can be
used to enhance bone and periodontal regeneration and aid in tissue
transports or bone grafts; prevents skin aging due to sunburn by
stimulating keratinocyte growth; prevents hair loss, since FGF family
members activate hair-forming cells and promotes melanocyte growth;
stimulates growth and differentiation of hematopoietic cells and bone
marrow cells when used in combination with other cytokines; maintains
organs before transplantation or for supporting cell culture of primary
tissues; induces tissue of mesodermal origin to differentiate in early
embryos; increases or decreases the differentiation or proliferation of
embryonic stem cells, besides, haematopoietic lineage; modulates
mamalian characteristics, such as, body height, weight, hair colour, eye
colour, skin, percentage of adipose tissue, pigmentation, size, and shape
(e.g., cosmetic surgery); modulates mammalian metabolism; changes
mammary's metal state or physical state by influencing biotryhms,
cardiac rhythms, depression, tendency for violence, tolerance for pain,
reproductive capabilities, hormonal or endocrine levels, appetite,
libido, memory, or stress; increases or decreases storage capabilities,
fat content, lipid, protein, carbohydrate, vitamins, minerals, cofactors
or other nutritional components. This sequence encodes a novel human
musculoskeletal system antigen.
Note: The sequence data for this patent did not form part of the
printed specification, but was obtained in electronic format directly
from the US patent office at
ftp.seqdata.uspto.gov/sequence.html?DocID=20020347140.
Sequence 9192 BP; 2180 A; 1723 C; 1845 G; 3444 T; 3 other;
Query March 86.7%; Score 434.4; DB 25; Length 9192;
Best Local Similarity 92.1%; Pred. No. 14e-107;
Matches 475; Conservative 0; Mismatches 26; Indels 15; Gaps 1;
QY 1 GGTGTTGGTGGAGTTCTTAATCTCGAGTCTCTAGTTTATTCAGTCTGGCTGAGAGAC 60
DB 6862 GGTGTTGGTGGAGTCTTAATCTCGAGTCTCTAGTTTATTCAGTCTGGCTGAGAGAT 6921
QY 61 AGTTGTTGTAATTTCTGTTCTTTACATCTGCTGAGGAGTGGTTAGTCTCACTATGT 120
DB 6922 AGTTGTTGTAATTTCTGTTCTTTACATCTGCTGAGGAGGCTTACTCTCAAGTATGT 6981
QY 121 GGTCAATTTGGAAATAGTGTGGTGGTCTGAGAGAGATGATATCTGTTGATTTGG 160
DB 6982 GGTCAATTTGGAAATAGTGTGGTGGTGGTCTGAGAGAGATGATATCTGTTGATTTGG 7041
QY 181 GGTGTTAGATTTCTGTAGATGTTCTATTAGTCTCACTTGGTGGAGAGTGGATTCAGTTCT 240
```

C Producing transgenic animals and cells and also in gene therapy. This
C sequence represents a prostate specific nucleic acid described in
C the invention.

X C Sequence 1885 BP; 662 A; 421 C; 395 G; 404 T; 0 other;

Query Match 85.4%; Score 432.8; DB 24; Length 1885;

Best Local Similarity 91.9%; Pred. No. 2.4e-107;

Matches 474; Conservative 0; Mismatches 27; Indels 15; Gaps 1;

Y 1 GGTATTGGTGGATGTTCTTAATCTGTAGTCTAGTTGATTTGACATGTGGCCCGAGAGAC 60

b 733 GGTATTGAGTGAGTTCTTAATCTGTAGTCTAGTTGATTTGACATGTGGCCCGAGAGAC 674

Y 61 AGTTTGTTGTAATTTCTGTTCTTTTATGTTGTTGAGGAGTGTTTATGTTCCAACTATGT 120

b 673 AGTTTGTTGTAATTTCTGTTCTTTTATGTTGTTGAGGAGTGTTTATGTTCCAACTATGT 614

Y 121 GGTCAATTTTGGATAGTGTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180

b 613 GGTCAATTTTGGATAGTGTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 554

Y 181 GGTATAGAGTCTCTAGATGTTCTATTTAGGTCACATTTGTCACAGAGTGTGATTCAGTCT 240

b 553 GGTGAGAGTCTCTAGATGTTCTATTTAGGTCACATTTGTCACAGAGTGTGATTCAGTCT 494

Y 241 CGA-----TCTGTCTTTGATCTGTCTATATATGACAGTGGGTTGTA 285

b 493 GGAATCTCTTTAACCTTTGTCATTTGATTTGATTTGATTTGATTTGATTTGATTTGAT 434

Y 286 AGTCTCCAGATTTATTTGTTGGAGTCTAAGTCTCTTTTATGTTCTTAGGAGTCTGT 345

b 433 AGTCTCCAGATTTATTTGTTGGAGTCTAAGTCTCTTTTATGTTCTTAGGAGTCTGT 374

Y 346 TATGAACTCTGGTCT 405

b 373 TATGAACTCTGGTCT 314

Y 456 GTGAATTTGACCTTTTATGATTTGATTTGATTTGATTTGATTTGATTTGATTTGATTT 465

b 313 GTGAATTTGACCTTTTATGATTTGATTTGATTTGATTTGATTTGATTTGATTTGATTT 254

Y 466 GTTAAAGTCTGTTTATGACAGAGTGGATTGCA 501

b 253 GTTAAAGTCTGTTTATGACAGAGTGGATTGCA 215

ESJUT 9

D AAZ35351/c

X AAZ35351 standard; DNA; 41599 BP.

C AAZ35351.

C 27-MAR-2000 (first entry)

C Cosmid including sequence spanning human chromosome 9p21.

C Chromosome 9p21; human; cancer; tumour; diagnosis; prognosis;

C methylthioadenosine phosphorylase; glioma; melanoma;

C primary lymphoid malignancy; non-small cell lung cancer;

C head and neck cancer; ovarian cancer; bladder cancer;

C chondrosarcoma; ds.

C Homo sapiens.

C WO9867634-A1.

C 29-DEC-1999.

C 21-JUN-1999; 99WO-051398.

C 23-JUN-1998; 98US-069041.

C 17-JUN-1999; 99US-0335231.

XX (REGC) UNIV CALIFORNIA.

XX Carson DA, Schmid M, Carrera CJ;

XX WPI: 2000-126650/11.

XX Diagnosing and determining prognosis for cancer causatively associated
PT with derangements of chromosome 9p21 -

XX Disclosure: Fig 7; 55pp; English.

XX This is the nucleotide sequence of a cosmid that includes sequences
CC spanning human chromosome 9p21 (AF0040047). This region
CC harbours 5 genes within about 120 kb: the tumour suppressor genes
CC p15INK4B (p15) with its alternatively spliced form p10, p16INK4A
CC (p16) and p14ARF, and the gene for methylthioadenosine phosphorylase
CC (MTAP). The invention provides a method for diagnosis of, and
CC determining a prognosis for, cancer causatively associated with
CC derangements of chromosome 9p21. Underlying the invention is the
CC discovery that such derangements have their genesis in deletions
CC occurring centromeric to STR 3.21, most often including breakpoints
CC in exon 5 and/or between exons 4 and 5 of the gene which encodes
CC MTAP. As the cancer and tumour development advance, deletions in
CC 9p21 progress centromerically from the genesis point toward the
CC gene encoding p16. Thus, the method of the invention is performed
CC by determining whether (a) portions of the 9p21 region including
CC and telomeric to STR 3.21 are deleted, and (b) portions of the
CC 9p21 region centromeric to STR 3.21 are deleted, where a positive
CC finding in step (a) and a negative finding in step (b) are
CC indicative of a cancer in an early stage of tumour development
CC and a positive finding in step (a) is indicative of a cancer in an
CC advanced stage of tumour development. Primer pairs (see BAZ35354-75)
CC are provided for use in claimed methods for diagnosing and
CC determining a prognosis for cancer associated with derangements of
CC 9p21, especially a glioma, primary lymphoid malignancy, non-small
CC cell lung cancer, melanoma, head and neck cancer, ovarian cancer,
CC bladder cancer or a chondrosarcoma (claimed).

XX Sequence 41599 BP; 13477 A; 8216 C; 8242 G; 11664 T; 0 other;

Query Match 86.1%; Score 431.6; DB 21; Length 41599;

Best Local Similarity 91.8%; Pred. No. 1.3e-106;

Matches 472; Conservative 0; Mismatches 29; Indels 13; Gaps 1;

Y 1 GGTATTGGTGGATGTTCTTAATCTGTAGTCTAGTTGATTTGACATGTGGCCCGAGAGAC 60

b 1096 GGTATTGAGTGAGTTCTTAATCTGTAGTCTAGTTGATTTGACATGTGGCCCGAGAGAC 11937

Y 61 AGTTTGTTGTAATTTCTGTTCTTTTATGATTTGCTGAGGAGTGTCTTAGTTCCAACTATGT 120

b 11834 AGTTTGTTGTAATTTCTGTTCTTTTATGATTTGCTGAGGAGTGTCTTAGTTCCAACTATGT 11777

Y 121 GGTCAATTTTGGAAATAGGTGTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 182

b 11776 GGTCAATTTTGGAAATAGGTGTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 11717

Y 181 GGTATAGAGTCTCTAGATGTTCTATTTAGGTCCTCACTGGTCCAGAGTGTAGTTCTCT 240

b 11716 GGTGGAGAGTCTCTAGATGTTCTATTTAGGTCCTCACTGGTCCAGAGTGTAGTTCTCT 11657

Y 241 CGA-----TCTGTCTTTGTTGATCTGTCTATATTTGACAGTGGGGTGTGGAAG 287

b 11656 GGATATCTCTTTAACTTTCTCTCACTGATCTCTCACTGATCTCTCACTGATCTCTCTTAAAG 11597

Y 288 TCTCCAGATTTATTTGTTGGGAGTCTTAAGTCTCTTTGTTAGGTCCTTAGGAGTCTCTCT 347

b 11536 TCTCCAGATTTATTTGTTGGGAGTCTTAAGTCTCTTTGTTAGGTCCTTAGGAGTCTCTCT 11537

Y 348 ATGAATCTGGGTGCTCTCTGTTATTTGGTGCATATATATTAGGATAGTTAGCTCTTCTGT 407

b 11536 ATGAATCTGGGTGCTCTCTGTTATTTGGTGCATATATATTAGGATAGTTAGCTCTTCTGT 11477

The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome mapping and gene mapping, and in recombinant production of (II). The polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (II) is useful in gene therapy techniques to restore normal activity of (I) or to treat disease states involving quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical imaging of sites expressing (II). (I) and (II) are useful for treating disorders involving aberrant protein expression or biological activity. The polypeptide and polynucleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic coding sequences of the invention.

Note: the sequence data for this patent did not appear in the printed specification, but was obtained in electronic format directly from WIPo at ftp.wipo.int/pub/published_pct_sequences.

Sequence 2606 BP; 982 A; 603 C; 478 G; 546 T; 0 other;

Query Match 86.1%; Score 431.2; DB 23; Length 2606;
Best Local Similarity 91.7%; Pred. No. 7.2e-10;
Matches 473; Conservative 0; Mismatches 28; Indels 15; Gaps 1

1 GGTITGGTGAGTTCCTAAATCGTAGCTCCTGATTGCACCTGGGCTGAGAGAC 60
1443 GGTTTGAGTGAGTTCCTTAATCGTAGCTCCTGATTGCACCTGGGCTGAGAGAC 1364
61 AGTTTGTTGAATTCCTCTTTTACATTCCTCGAGAGAGCTTTAGTTCGACTATGT 120
1383 AGTTTGTTGAATTCCTCTTTTACATTCCTCGAGAGAGCTTTAGTTCGACTATGT 1324
121 GGTCAATTTGGAATAGGCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
1323 GGTCAATTTGGAATAGGCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 1264
131 GTTAGAGTTCCTGAGTATGTTTATAGTCCACTGGTGGAGAGCTGAGTTCAGTTTCT 240
1263 GGTGGAGAGTTCCTGAGTATGTTTATAGTCCACTGGTGGAGAGCTGAGTTCAGTTTCT 1264
241 GGA-----TCTGT 283
1253 GGATATCCTGTGTACCTTCTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGT 1144
286 AGTCTCCCAGTATTTGTTGGGAGCTTAAGTCTCTTTGAGGCTCTTATAGGAGCTTCT 345
1143 AGTCTCCCAGTATTTGTTGGGAGCTTAAGTCTCTTTGAGGCTCTTATAGGAGCTTCT 1084
346 TTATCAATCTGGTCTCTGTATGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 405
1083 TTATCAATCTGGTCTCTGTATGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 1024
406 GTTGAATTCATCCCTTTACCATTTATATGATGCCCTCTTTGTCTTTGTGATCTTTGTG 465
1023 GTTGAATTCATCCCTTTACCATTTATATGATGCCCTCTTTGTCTTTGTGATCTTTGTG 564
466 GTTGAATTCATCCCTTTACCATTTATATGATGCCCTCTTTGTCTTTGTGATCTTTGTG 502
963 GTTGAATTCATCCCTTTATCAGAGAGCTAGGATTCGAA 928

RESULT 12
D AAS70259 standard; cDNA; 3316 BP.

11-OCT-2001; 2001WO-US08631.
 30-MAR-2001; 2001WO-US08631.
 31-MAR-2000; 2000US-5640217.
 23-AUG-2000; 2000US-5649167.
 (HYSB-) HYSB INC.
 Drmanac RT, Liu C, Tang YT;
 WPI; 2001-639362/73.
 P-PSDB; ABG10813.
 New isolated polynucleotide and encoded polypeptides, useful in
 diagnostics, forensics, gene mapping, identification of mutations
 responsible for genetic disorders or other traits and to assess
 biodiversity -
 Claim 1: SEQ ID NO 9644; 103pp; English.
 The invention relates to isolated polynucleotide (I) and
 polypeptide (II) sequences. (I) is useful as hybridisation probes,
 polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 and gene mapping, and in recombinant production of (II). The
 polynucleotides are also used in diagnostics as expressed sequence tags
 for identifying expressed genes. (I) is useful in gene therapy techniques
 to restore normal activity of (II) or to treat disease states involving
 quantitating a polypeptide in tissue, as molecular weight markers and as
 a food supplement. (II) and its binding partners are useful in medical
 imaging of sites expressing (II). (I) and (II) are useful for treating
 disorders involving aberrant protein expression or biological activity.
 The polypeptide and polynucleotide sequences have applications in
 diagnostics, forensics, gene mapping, identification of mutations
 responsible for genetic disorders or other traits to assess biodiversity
 and to produce other types of data and products dependent on DNA and
 amino acid sequences. AAS64157-AAS94564 represent novel human
 diagnostic coding sequences of the invention.
 Note: The sequence data for this patent did not appear in the printed
 specification, but was obtained in electronic format directly from WIPCO
 at ftp.wipo.int/pub/published_pct_sequences.
 Sequence 3316 BP: 1389 A; 713 C; 561 G; 653 T; 0 other.
 Query March 96.1% Score 431.2; DB 23; Length 3316;
 Best Local Similarity 91.7% Pred No 7.8e-107;
 Matches 473; Conservative 0; Mismatches 28; Indels 15; Gaps 1;
 1 GGTTCGGTGAGTCTTTAAAGCTGAGTCTAGTTTGAATTCAGTCTGAGTCTGAGAGAC 60
 729 GGTTCGGTGAGTCTTTAAAGCTGAGTCTAGTTTGAATTCAGTCTGAGTCTGAGAGAC 672
 61 AGTTGTTGAATTCGTTCTTTTACATTCGTTGAGGAGTCTGTTTACGAGTCTG 123
 669 AGTTGTTGAATTCGTTCTTTTACATTCGTTGAGGAGTCTGTTTACGAGTCTG 610
 121 GGTCAATTTGGGAATAGGTGGTGGTGGTCTGAGAGAGATGATATTCGTTGTTGG 180
 609 GGTCAATTTGGGAATAGGTGGTGGTGGTCTGAGAGAGATGATATTCGTTGTTGG 550
 181 GGTTCAGATCTGTAGATCTTATAGTCCACTTGGTGCAGAGCTGAGTCACTTCT 240
 549 GGTTCAGATCTGTAGATCTTATAGTCCACTTGGTGCAGAGCTGAGTCACTTCT 490
 241 GGA-----TCTGCTCTGATCTGCTCTATATTCAGATCGGGCTTGA 265
 489 GGTATCTCTGTTAACTCTCTCTCTGCTGATCTGCTAAATGTTGACGTTGGTTAA 430
 286 AGTCTCCAGTATATTGTGTGGAGTCTAAGTCTCTTTGTAGGCTCTTAGGAGCTTGT 345
 429 AGTCTCCAGTATATTGTGTGGAGTCTAAGTCTCTTTGTAGGCTCTTAGGAGCTTGT 370

QY 346 TTATGAATCTGGTCTCTCTGTTATTTGGTGTGATATATATTAGGATAGTCTCTCTT 405
 DB 363 TTATGAATCTGGTCTCTCTGTTATTTGGTGTGATATATATTAGGATAGTCTCTT 310
 QY 436 GTTGATTCGATCCCTTAGCA-TATATGATGCGCTTCTTTCTCTTTTGTATCTTTG 465
 DB 309 GTTGATTCGATCCCTTTACCATCATGTAAAGGCCCTTTCTCTCTTTGATCTTTG 250
 QY 466 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 501
 DB 249 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 214
 RESULT 15
 AAS7500C/C
 C AAS7500C standard; cDNA; 3316 BP.
 AC AAS7500C;
 DT 13-FEB-2002 (first entry)
 XX DNA encoding novel human diagnostic protein #13804.
 DE Human; chromosome mapping; gene mapping; gene therapy; forensics;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 OS Homo sapiens.
 XX W0200175067-A2.
 XX 11-OCT-2001.
 PF 30-MAR-2001; 2001WO-US08631.
 PR 31-MAR-2000; 2000US-5640217.
 PR 23-AUG-2000; 2000US-5649167.
 XX (HYSB-) HYSB INC.
 XX Drmanac RT, Liu C, Tang YT;
 XX WPI; 2001-639362/73.
 XX P-PSDB; ABG10813.
 XX New isolated polynucleotide and encoded polypeptides, useful in
 diagnostics, forensics, gene mapping, identification of mutations
 responsible for genetic disorders or other traits and to assess
 biodiversity -
 Claim 1: SEQ ID NO 13804; 103pp; English.
 The invention relates to isolated polynucleotide (I) and
 polypeptide (II) sequences. (I) is useful as hybridisation probes,
 polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 and gene mapping, and in recombinant production of (II). The
 polynucleotides are also used in diagnostics as expressed sequence tags
 for identifying expressed genes. (I) is useful in gene therapy techniques
 to restore normal activity of (II) or to treat disease states involving
 quantitating a polypeptide in tissue, as molecular weight markers and as
 a food supplement. (II) and its binding partners are useful in medical
 imaging of sites expressing (II). (I) and (II) are useful for treating
 disorders involving aberrant protein expression or biological activity.
 The polypeptide and polynucleotide sequences have applications in
 diagnostics, forensics, gene mapping, identification of mutations
 responsible for genetic disorders or other traits to assess biodiversity
 and to produce other types of data and products dependent on DNA and
 amino acid sequences. AAS64157-AAS94564 represent novel human
 diagnostic coding sequences of the invention.
 Note: The sequence data for this patent did not appear in the printed
 specification, but was obtained in electronic format directly from WIPCO
 at ftp.wipo.int/pub/published_pct_sequences.


```
SC Sequence 3316 BP; 1389 A; 713 C; 561 G; 653 T; 0 other;
Query Match 96.1%; Score 431.2; DB 23; Length 3316;
Best Local Similarity 91.7%; Pred. No. 7.9e-107;
Matches 473; Conservative 0; Mismatches 28; Indels 15; Gaps 1;
QY 1 GGTTCGCGTGGAGTTCCTTAATCGAGTCTAGTTCGATTCGACGTGSGCTGAGAGAC 60
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
729 GGTTCGAGTGGAGTTCCTTAATCGAGTTCGAGTTCGAGTTCGAGTTCGAGAGAC 170
QY 61 AGTTGTGTGAATTCCTGTTCCTTTACATTCGTGAGAGTTCCTTAGTTCGAACTATGT 120
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 669 AGTTGTGTGAATTCCTGTTCCTTTTATATTCGTGAGAGTTCCTTAGTTCGAACTATGT 410
QY 121 GGTCAATTTTGGAAATAGGTGTGTGTGCTGAGAGAGATGATATTCGTGTGATTTGG 180
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 609 GGTCAATTTTGGAAATAGGTGTGTGTGCTGAGAGAGATGATATTCGTGTGATTTGG 450
QY 181 GGTTCGAGTTCGTGATGATGCTATTCAGTCCACTTGGTGCAGAGCTGAGTTCAGTTGCT 240
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 549 GGTGGAGAGTTCGTGATGATGCTATTCAGTCCACTTGGTGCAGAGCTGAGTTCAGTTGCT 490
QY 241 GGA-----CTGTCTCTGTTCATCTGTCTTAATTCAGAGTGGAGTCTTGA 485
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 489 GGATATCCTTGTTAACTTCCTGTCTCTGTTCATCTGTCTTAATTCAGAGTGGAGTCTTGA 430
QY 286 AGTCCCGCAATATATGTGTGGAGTCTAAGTCTCTTTGTAGTCTCTAGGAGCTTGCT 345
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 429 AGTCCCGCAATATATGTGTGGAGTCTAAGTCTCTTTGTAGTCTCTAGGAGCTTGCT 370
QY 346 TTATGAATCTGGGTGCTCCTGTATTGGTGCACATATATTAGGATAGTTCGTCTCTT 405
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 369 TTATGAATCTGGGTGCTCCTGTATTGGTGCACATATATTAGGATAGTTCGTCTCTT 310
QY 406 GTTGAATTCACCTTTAGCAATTAATATGATGGCTTCCTTTCTCTTTGATCTTTTGTG 465
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 309 GTTGAATTCACCTTTAGCAATCATGTAATGGCTTCCTTTCTCTTTGATCTTTTGTG 250
QY 466 GTTAAAGTCTGTTTATCAGAGATTCGATTCGAA 501
CB ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 249 GTTAAAGTCCGTTTATCAGAGACTAGGATTCGAA 214
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Search completed: October 24, 2003, 18:47:22
Job time : 131.251 secs


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52128 AGCTTCACATATTAAATGCTGGAGCTAAGTCTCTTTTAGTCTCACTGAGGACTTGGT 52187
346 TTATGAATCTGGAGCTCTCTATGCTGGTGCATATATATTAGGATAGTCTTCTT 405
52188 TTATGAATCTGGAGCTCTCTATGCTGGTGCATATATATTAGGATAGTCTTCTT 52247
406 GTTGAATGATCCCTTACGATATATAGTGGCTCTTCTCTCTTTAGTCTTCTT 465
52248 GTTGAATGATCCCTTACGATATATAGTGGCTCTTCTCTCTTTAGTCTTCTT 52307
466 GTTAAAGTCTGTTTATCAGAGAGTTCGAA 501
52308 GTTAAAGTCTGTTTATCAGAGAGTTCGAA 52343

RESULT 2
US-09-426-290-1
; Sequence 1, Application US/09426290
; Patent No. 6410912
; GENERAL INFORMATION:
; APPLICANT: Berglund Rn Olafsdottir
; APPLICANT: Jeffrey Guicher
; TITLE OF INVENTION: HUMAN NARCOLEPSY GENE
; FILE REFERENCE: 2345.2001-000
; CURRENT APPLICATION NUMBER: US/09/426.290
; CURRENT FILING DATE: 1999-10-25
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: PastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 168575
; TYPE: DNA
; ORGANISM: Homo Sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (21181)...(121403)
; NAME/KEY: CDS
; LOCATION: (95252)...(195430)
; NAME/KEY: CDS
; LOCATION: (101753)...(101996)
; NAME/KEY: CDS
; LOCATION: (110324)...(110439)
; NAME/KEY: CDS
; LOCATION: (124058)...(124278)
; NAME/KEY: CDS
; LOCATION: (127009)...(127130)
; NAME/KEY: CDS
; LOCATION: (128910)...(129139)
; US-09-426-290-1

Query Match 84.9%; Score 425.4; DB 4; Length 168575;
Best Local Similarity 91.1%; Pred. No. 4.7e-113;
Matches 469; Conservative 0; Mismatches 32; Indels 15; Gaps 1;

QY 2 GTTTTCGGTGAGTTTCTTAATCCTCAGTTCTAGTTGATGCTGAGGAGTCTTCTT 156093
DB 156093 GTTTTCGGTGAGTTTCTTAATCCTCAGTTCTAGTTGATGCTGAGGAGTCTTCTT 156152
QY 62 GTTTTCGTTAATTTCTGTTCTTTTACATTTGCTGAGGAGTCTTCTTCTTCTTCTT 156153
DB 156153 GTTTTCGTTAATTTCTGTTCTTTTACATTTGCTGAGGAGTCTTCTTCTTCTTCTT 156212
QY 122 GTCAATTTTGGAAATAGTGTGGTGGTCTCAGAGAAATGTATATTCTGTTGATTGGG 181
DB 156213 GTCAATTTTGGAAATAGTGTGGTGGTCTCAGAGAAATGTATATTCTGTTGATTGGG 156272
QY 182 GTTTAGAGTTCTGTAGATGCTATATTAGTCCACTTGGTGCAGAGTCTGAGTTCTCTG 241
DB 156273 GTTGAGAGTTCTGTAGATGCTATATTAGTCTCTTCTGTCAGAGTCTGAGTTCTCTG 156332
QY 242 GA-----TCTGCTCTGTTGATCTGTCTAATATTGACAGTGGGGTGTGAA 296
DB 156333 GATATCCTGTAACTTTCTGCTGATCTGTTATGTTGACAGTGGGGTGTGAA 156392
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QY 287 GTCTCCACGATATATTGCTGGAGTCTAAGTCTCTTTTAGTCTCTTAGGACTTGGT 346
DB 156193 GTCTCCACGATATATTGCTGGAGTCTAAGTCTCTTTTAGTCTCTTAGGACTTGGT 156452
QY 347 TATGAATCTGGGTCCTCTGTTATGGTGCATATATATTAGGATAGTCTTCTTCTT 406
DB 156453 TATGAATCTGGGTCCTCTGTTATGGTGCATATATATTAGGATAGTCTTCTTCTT 156512
QY 407 TTGAATGATCCCTTACGATATATAGTGGCTCTTCTTCTCTTTAGTCTTCTTCTT 466
DB 156513 TTGAATGATCCCTTACGATATATAGTGGCTCTTCTTCTCTTTAGTCTTCTTCTT 156572
QY 466 TTGAAGTCTGTTTATCAGAGAGTTCGAA 501
DB 156573 TTGAAGTCTGTTTATCAGAGAGTTCGAA 156607

RESULT 3
US-09-245-281-44
; Sequence 44, Application US/09245281
; Patent No. 6365196
; GENERAL INFORMATION:
; APPLICANT: Bestin, John
; TITLE OF INVENTION: NOVEL MOLECULES OF THE CARD-RELATED PROTEIN FAMILY
; FILE REFERENCE: 07334/11800
; CURRENT APPLICATION NUMBER: US/09/245.281
; CURRENT FILING DATE: 1999-02-25
; EARLIER APPLICATION NUMBER: US 09/207,359
; EARLIER FILING DATE: 1998-12-08
; EARLIER APPLICATION NUMBER: US 09/099,041
; EARLIER FILING DATE: 1998-06-17
; EARLIER APPLICATION NUMBER: US 09/019,942
; EARLIER FILING DATE: 1998-02-06
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PastSeq for Windows Version 4.0
; SEQ ID NO 44
; LENGTH: 32042
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-245-281-44

Query Match 84.8%; Score 424.8; DB 4; Length 32042;
Best Local Similarity 90.9%; Pred. No. 3.9e-113;
Matches 469; Conservative 0; Mismatches 32; Indels 15; Gaps 1;

QY 1 GGTTCGCTGAGTTTCTTAATCCTCAGTTCTAGTTGATGCTGACCTGTGGCTGAGAGAC 60
DB 15442 GGTTCGCTGAGTTTCTTAATCCTCAGTTCTAGTTGATGCTGACCTGTGGCTGAGAGAT 15501
QY 61 AGTTTCGTTAATTTCTGTTCTTTTACATTTGCTGAGGAGTCTTCTTCTTCTTCTTCTT 15502
DB 15502 AGTTTCGTTAATTTCTGTTCTTTTACATTTGCTGAGGAGTCTTCTTCTTCTTCTTCTT 15561
QY 121 GGTCAATTTTGAATAGTGTGGTGGTCTGAGAGAAATGTATATTCTGTTGATTTGG 180
DB 15562 GGTCAATTTTGAATAGTGTGGTGGTCTGAGAGAAATGTATATTCTGTTGATTTGG 15621
QY 181 GGTTCGAGTTCTGTAGATGCTATATTAGTCCACTTGGTGCAGAGTCTGAGTTCTCTT 240
DB 15622 GGTTCGAGTTCTGTAGATGCTATATTAGTCTCTTGGTGCAGAGTCTGAGTTCTCTTCT 15681
QY 241 GG-----ATCTGCTCTTGTGATCTGCTAATATTGACAGTGGGGTCTTGA 285
DB 15682 GGATATCTCTTGTGATCTGCTAATATTGACAGTGGGGTCTTGA 15741
QY 286 AGTCTCCAGTATTATTTGTTGGAGTCTAAGTCTCTTTTAGTCTCTTAGGACTTGGT 345
DB 15742 AGTCTCCAGTATTATTTGTTGGAGTCTAAGTCTCTTTTAGTCTCTTAGGACTTGGT 15801
QY 346 TTATGAATCTGGTGGTCTCTGTATTTAGGTCATATATTAGGATAGTCTTCTTCTTCTT 405
DB 15802 TTATGAATCTGGTGGTCTCTGTATTTAGGTCATATATTAGGATAGTCTTCTTCTTCTT 15861
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; SEQ ID NO 8
; LENGTH: 18597
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: 701..1375;
; OTHER INFORMATION: n = c or a
; NAME/KEY: misc feature
; LOCATION: 716..1293; 2401..2429; 2518..3083; 3125..3435; 4254..4898;
; LOCATION: 5062..5167; 11069..13299; 14479..14730; 14796..15344; 15450;
; OTHER INFORMATION: n = a or g
; NAME/KEY: misc feature
; LOCATION: 732..1379; 1597..2488; 3212..5066; 11238..11422; 11666;
; LOCATION: 12598; 13171..13645; 13782..13826; 13913..14586; 14788;
; LOCATION: 15042..15546; 15770
; OTHER INFORMATION: n = c or t
; NAME/KEY: misc feature
; LOCATION: 1322..1688
; OTHER INFORMATION: n = c or g
; NAME/KEY: misc feature
; LOCATION: 2594..11293; 16199..16203
; OTHER INFORMATION: n = g or t
; NAME/KEY: misc feature
; LOCATION: 3619
; OTHER INFORMATION: n = a or t
; NAME/KEY: misc feature
; LOCATION: 14547
; OTHER INFORMATION: nucleotide in position 14547 is t, or absent
;S-09-962-665-8

Query Match 45.9%; Score 230; DB 4; Length 18597;
Best Local Similarity 87.4%; Pred. No. 6.9e-57;
Matches 304; Conservative 0; Mismatches 25; Indels 19; Gaps 4;

Y 2 GTTTTCGCGAGTTTCTTAATCCTGAGTTCTAGTTGATTCAGTCTGGCTGAGAGACA 61
b 9059 GTTTTGGAGTCTTCAATCCTGAGTTCTAGTTGATTCAGTCTGGCTGAGAGACA 9113
Y 62 GTTTTCGCGAGTTTCTTAATCCTGAGTTCTAGTTGATTCAGTCTGGCTGAGAGACA 121
b 9113 GTTTTGGAGTCTTCAATCCTGAGTTCTAGTTGATTCAGTCTGGCTGAGAGACA 9178
Y 122 GTCAATTTGGAAATAGGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 181
b 9178 GTTCGTTTGGAAATAGGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 9238
Y 182 GTTTAGAGTTCTGTAGATCTCTATATTAGTCTGCTGCTGCTGCTGCTGCTGCTG 241
b 9239 GTTTCGTTTGGAAATAGGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 9296
Y 242 G-----ATCGTCTTGTGATCTGCTATATTGACAGTGGGTGGTGGAA 286
b 9297 GTTATCCTTGTGACTTTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 9355
Y 287 GTCTCCAGTATTATTGTGGAGTCTAAAGTCTCTTTGTAGTCTCT 334
b 9356 GTCTCCAGTATTATTGTGGAGTCTAAAGTCTCTTTGTAGTCTCT 3402

RESULT 13
S-09-525-160B-9/c
Sequence 9, Application US/59525160B
Patent No. 6569681
GENERAL INFORMATION:
APPLICANT: Ivanov, Evgenii
TITLE OF INVENTION: METHODS OF IMPROVING HOMOLOGOUS RECOMBINATION
FILE REFERENCE: 10278/016001
CURRENT APPLICATION NUMBER: US/09/525,160B
CURRENT FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 10
SOFTWARE: FastSeq for Windows Version 4.0

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; SEQ ID NO 9
; LENGTH: 3033
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-525-160B-9

Query Match 38.4%; Score 192.2; DB 4; Length 3033;
Best Local Similarity 65.5%; Pred. No. 3.1e-46;
Matches 319; Conservative 0; Mismatches 153; Indels 15; Gaps 2;

QY 18 TTAATCCTCAGTTCTAGTTTGTGATTCAGTCTGGCTGAGAGACAGTTTGTGTAATTCT 77
b 2074 TTATTGGGACTCTCTTTTCTTCTAGTCTGTGGTGGAGAGAAAGTTGATATTCTCA 2015
QY 78 GTCTCTTTACATTTCTCTGAGGAGTCTTGTAGTTCTCACTATGTTGTAATTTTGAATAG 137
b 2014 CTTTTCTTCTAAATTTTAAAGATTTGTTTGTAGCTTGTAGCTTGTATATTCCTTGA 1955
QY 138 GTGTGGTGTGGTGTGAGAGAAATGTAATTTCTGTGATTTGGGGTTTAGAGTTCTCTAG 197
b 1954 TATCCATGTGTGAGGAGAGAAATGTGATTTCTAGAGTGTGATCAATGTTCTGTAA 1895
QY 198 ATGTCTATTAGTCCACTTGTGGCAGAGCTGAGTTCA-----GTTCCTGGATCT 246
b 1894 ATATCTACTAGTCCACTTGTAGCTTGTAGCTTGTAGCTTGTAGCTTGTAGCTTGTAG 1835
QY 247 GTCTTGTGAGTCTGTCTAATATTGACAGTGGGTGTGAGCTTCCAGTATTTATTTGT 306
b 1834 GTCTGATGATCTGTGCAATCTGAAAGTGGGTGTGAAAGTCCAGTATTTATTTAT 1775
QY 307 GGG-----AGTCTAACTCTCTTTGTAGTCTCTAGGAGTCTCTTATGAACTGGGTCT 362
b 1774 GGGGGTCTGTCTATCTCTCTTTGGCTCTAACAATATTGCTTTATATACCTGAGTCT 1715
QY 363 CTGTATTTGGTGGCATATATTAGGATAGTCTGCTTCTTGTGTAATTTGATCCCTTT 422
b 1714 CCAGTATTTGGTGGTAAATATATTAGAAATTTGTAATCTCTTCTTCTTCTTCTTCT 1655
QY 423 ASCATTATATGATGGCTCTCTTTGTCTCTTTGTCTTTGTCTTTGTCTTTGTCTTTGT 482
b 1654 ATCATATATAAGACCTCTCTTCTCTCTTCTCTCTCTCTCTCTCTCTCTCTCTCT 1595
QY 483 TCAGAGA 489
b 1594 TCTGATA 1588

RESULT 14
US-09-525-160B-4/c
Sequence 4, Application US/09525160B
Patent No. 6569661
GENERAL INFORMATION:
APPLICANT: Ivanov, Evgenii
TITLE OF INVENTION: METHODS OF IMPROVING HOMOLOGOUS RECOMBINATION
FILE REFERENCE: 10279/016001
CURRENT APPLICATION NUMBER: US/09/525,160B
CURRENT FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 10
SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 4
; LENGTH: 3213
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-525-160B-4

Query Match 38.4%; Score 192.2; DB 4; Length 3213;
Best Local Similarity 65.5%; Pred. No. 3.1e-46;
Matches 319; Conservative 0; Mismatches 153; Indels 15; Gaps 2;

QY 18 TTAATCCTCAGTTCTAGTTTGTGATTCAGTCTGGCTGAGAGACAGTTTGTGTAATTCT 77
b 2074 TTATTGGGACTCTCTTTTCTTCTAGTCTGTGGTGGAGAGAAAGTTGATATTCTCA 2015

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QY 78 GTCTCTTTACATTTGCTGAGAGTGGTTTGTGTCACACTATGTGTCATATTTGGAAATAG 137
DB 2014 CTTCTTTCTTAAATTTTAAAGATTTGTTTGTGACCTAGCAATTTGGTATATATCTTGAAGT 1955
QY 138 GTGTGGTGTGGTGTGAGAGAAATGTAATTCCTGTGATTTGGGGTTTAGAGTCTCTGTAG 197
DB 1954 TATCATGTGCTGAGAGAGAGAAATGTGATTTCTACAGCTGTGAGTCAAAATGTTCTGTAA 1895
QY 198 ATGTCTATTAGTCCACTTTGTTGGCAGAGCTGAGTTCA-----GTTCCCTGGATCT 246
DB 1894 ATATCTACTAGGTCCATTTGACCTATGCTGTCAGATGTCAGATGAAATTTGAATTTTCT 1815
QY 247 GTCTGTTGATCTGCTATATATGACATGGGGTGTGAGTCTCCAGTCTCAATTTATGTGT 106
DB 1834 GTCTGGATGATCTGCCAATGCTGAAATGGGGTGTGAGTCTCCAGTCTCAATTTATGTGT 1775
QY 307 GGG-----AGCTTAAGTCTCTTTTATAGGCTCTGAGGACTTGGTTTATGAATCTGGTCT 362
DB 1774 GGGGGTCTGCTATCTCTCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 1715
QY 363 CTGTGATTGGTGCATATATATATAGGATAGTATAGTCTCTCTGTTGAACTGATCCCTTT 422
DB 1714 CAGTATTGGTGAATATATATATAGAACTGTTAAATGCTTTGTAATGGACCCCTTT 1655
QY 423 AGCATATCATGATGGCTCTCTCTGCTCTCTTTGATGCTTTGTTGTTGTTAAAGCTGTTTA 482
DB 1654 ATCACTATATATGACCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1595
QY 483 TCAGAGA 489
DB 1594 TCTGATA 1598

RESULT 15
US-09-220-132-168/c
; Sequence 168; Application US/09220:132
; Patent No. 6506607
; GENERAL INFORMATION:
; APPLICANT: Shyjan, Andrew W.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE IDENTIFICATION AND ASSESSMENT
; FILE REFERENCE: 07334-074001
; CURRENT APPLICATION NUMBER: US/09/220,132
; CURRENT FILING DATE: 1998-12-23
; PRIOR APPLICATION NUMBER: US 60/579,303
; PRIOR FILING DATE: 1998-03-25
; PRIOR APPLICATION NUMBER: US 60/568,821
; PRIOR FILING DATE: 1997-12-24
; NUMBER OF SEQ ID NOS: 191
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 168
; LENGTH: 9573
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-220-132-168

Query Match 37.4%; Score 187.4; DP 4; Length 9573;
Best Local Similarity 62.5%; Pred. No. 1.1e-44;
Matches 3.8; Conservative 0; Mismatches 176; Indels 15; Gaps 1;

QY 2 GTTTTGGCTGAGTTCTTAACTCTGAGTTCTAGTTTCACTGCTGAGCTTGGCTGAGAGACA 61
DB 4074 GTTTTAAAGGCTCTTTTGGAGTTAATTTCCASATTTTATTCGATTTGGTCTGMAAGCT 4015
QY 62 GTTTGTTGTAATTTCTGTTCTTTTACATTTGCTGAGAGAGTGTCTTGTAGTTCGAACTATGTG 121
DB 4014 ACTTGACATAATTTCAATTTTCTTAAATTTGTTGAGACTTGTCTGTGAGCTATCTTATG 3955
QY 122 GTCATTTTGGATAGGTGGTGTGGTGTGAGAGAACTGTAATTTCTGTTGATTTGG 161
DB 3954 ATTTATCTTGAGAGACCTTCTA-TGTGCTGATTAAATA-TGGTATATTTCTGAAAT-TGTGG 3695
QY 182 GTTTAGAGTTCTGTAGATGTCTATTAGTCCACTTGGTGCAGAGCTGAGTTCAGTTC... 238
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DB 3894 GTAGAACTGTAGTAAATATCTGCTAAATCTATTTTCTAGGATATAATTAAAGTCAAT 3835
QY 239 -----CTGGATCTGTCTTTGTTGACCTGTCTAATATTGACAGTGGGGTGTGGAA 286
DB 3834 TTTTCTTTATTGACATTTCTGCTTGTATGTCCTGTCTAGTCTGTGCCATGAGTATTGAA 1775
QY 287 GTTCCACAGTATTAATGTGTGGAGTCTAAGTCTCTTTGTAGTCTCTTAGGACTTGTCT 346
DB 3774 GTCCCTACTATTATTSTGTTGGTCCATCTATCCCATTTCTTAGGCTCTAGTAAATAATGTTT 3715
QY 347 TATGAATCTGGGTGCTCCTGTATCTGGGTSCATATATATTAGGATAGTTAGTCTCTCTTG 406
DB 3714 TATAAATTTGGGAGCTCCAGTATTAGTSCATATATATTAGGATTTGTATATTGTCTG 3655
QY 407 TTGAATTGATCCCTTTAGCATTATATGATGAGGCTCTCTTGTCTCTTTTGTATCTTTGTTG 466
DB 3654 TTGGACCGATCCCTTTATCATTAATAAAATTTCTCTCTTGTCTTTTATAAATGTTTGC 3595
QY 467 TTAAAGCTCTGTTTATCAGAGAGTTGGA 495
DB 3594 TTAAAGCTTGTCTGATAATAAAATAGGA 3566
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Search completed: October 24, 2003, 15:42:57
Cpu time : 36.3542 secs

GenCore version 3.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

DM nucleic - nucleic search, using sw model

Run on: October 24, 2003, 15:28:54 : Search time 123.415 Seconds
(without alignment)
10486.662 Million cell updates/sec

Title: US-09-830-902-1_COPY_50000_50500

Perfect score: 501

Sequence: 1 ggttttgctgagttctta.....accagagctggtattccaa 501

Scoring table: IDENTITY_NJC

Gapop 10.0, Gapext 1.0

Searched: 1792395 seqs, 134090345: residues

Total number of hits satisfying chosen parameters: 3534793

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Published Applications NA:
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2: /cgn2_6/ptodata/2/pubna/PCR_NEW_PUB.seq:
3: /cgn2_6/ptodata/2/pubna/US05_NEW_PUB.seq:
4: /cgn2_6/ptodata/2/pubna/US06_PUBCOMB.seq:
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12: /cgn2_6/ptodata/2/pubna/US08_NEW_PUB.seq:
13: /cgn2_6/ptodata/2/pubna/US09_PUBCOMB.seq:
14: /cgn2_6/ptodata/2/pubna/US10_PUBCOMB.seq:
15: /cgn2_6/ptodata/2/pubna/US11_NEW_PUB.seq:
16: /cgn2_6/ptodata/2/pubna/US12_NEW_PUB.seq:
17: /cgn2_6/ptodata/2/pubna/US13_PUBCOMB.seq:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	436	87.0	5065	14	US-10-050-982-39
2	436	87.0	1691139	14	US-10-067-524-1
3	434.4	86.7	9192	10	US-09-764-847-1246
4	434.4	86.7	9192	10	US-09-764-877-2218
5	434.4	86.7	9192	14	US-10-092-154-1246
6	431.2	86.1	5676	10	US-09-764-877-3756
7	431.2	86.1	32199	10	US-09-764-855-210
8	431.2	86.1	32199	14	US-10-072-349-220
9	431.2	86.1	250000	12	US-10-025-813-26
10	429.6	85.7	5446	11	US-09-764-891-7392
11	429.6	85.7	5671	10	US-09-764-877-3755
12	429.6	85.7	20746	11	US-09-764-891-10041
13	428	85.4	116592	10	US-09-848-512-3
14	428	85.4	116592	12	US-10-354-085-3
15	428	85.4	392000	12	US-10-027-983-11
16	428	85.4	465237	10	US-09-933-267A-1

17	428	85.4	786431	12	US-10-612-227-3
18	426.4	85.1	3591	9	US-09-764-869-1634
19	426.4	85.1	2693	14	US-10-091-524-1634
20	426.4	85.1	6181	10	US-09-764-864-1674
21	426.4	85.1	6539	14	US-10-025-201-1
22	426.4	85.1	6766	10	US-09-764-847-1678
23	426.4	85.1	6766	14	US-10-092-154-1678
24	425.4	84.3	168578	12	US-10-178-184-1
25	424.8	84.8	4779	11	US-09-764-891-6770
26	424.8	84.8	4779	14	US-10-091-572-6170
27	424.8	84.8	7384	11	US-09-764-891-6771
28	424.8	84.8	7384	14	US-10-091-572-6171
29	424.8	84.8	30515	10	US-09-764-847-1208
30	424.8	84.8	30515	14	US-10-092-154-1208
31	424.8	84.8	32042	9	US-09-728-721-63
32	424.8	84.8	32042	13	US-10-118-984-44
33	424.8	84.8	32042	14	US-10-295-981-63
34	424.8	84.8	34001	14	US-10-006-883A-15
35	424.8	84.8	41104	9	US-09-816-685-3
36	424.8	84.8	180557	13	US-10-003-806-6
37	424.8	84.8	180557	13	US-10-003-806-9
38	424	84.6	24132	9	US-09-764-860-661
39	424	84.6	24132	14	US-10-074-095-661
40	423.8	84.6	108359	14	US-10-191-807-3
41	423.2	84.5	7380	11	US-09-764-891-6772
42	423.2	84.5	7380	14	US-10-091-572-6172
43	423.2	84.5	8041	11	US-09-764-891-6738
44	423.2	84.5	8041	11	US-09-764-891-6740
45	423.2	84.5	8041	14	US-10-091-572-588

ALIGNMENTS

RESULT :

US-10-350-982-39
: Sequence 39, Application US/10350882
: Publication No. US2003010406A;
: GENERAL INFORMATION:
: APPLICANT: Ruben et al.
: TITLE OF INVENTION: 27 Human secreted proteins
: FILE REFERENCE: P2038P1
: CURRENT APPLICATION NUMBER: US/10350,882
: CURRENT FILING DATE: 2002-02-18
: PRIOR APPLICATION NUMBER: 09/661,453
: PRIOR FILING DATE: 2000-09-13
: PRIOR APPLICATION NUMBER: PCT/US00/06793
: PRIOR FILING DATE: 2000-03-16
: PRIOR APPLICATION NUMBER: 607,125,055
: PRIOR FILING DATE: 1999-03-18
: NUMBER OF SEQ ID NOS: 156
: SOFTWARE: Patent'n Ver. 2.0
: SEQ ID NO 39
: LENGTH: 5065
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: SITE
: LOCATION: 12531;
: OTHER INFORMATION: n equals a.t.g. or c
: NAME/KEY: SITE
: LOCATION: 15063;
: OTHER INFORMATION: n equals a.t.g. or c
US-10-350-982-39

Query Match 87.0%; Score 436; DB 14; Length 5065;
Best Local Similarity 91.7%; Pred. No. 1.3e-108;
Matches 473; Conservative 4; Mismatches 24; Indels 15; Gaps 1;

Oy 1 GGTTCGGTGGATTCCTTAACTCGTGGTCTAGTTGATTCGCTGGCTGAGAGAC 60
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 3355 GGTTCGGTGGATTCCTTAACTCGTGGTCTAGTTGATTCGCTGGCTGAGAGAY 3414

405 GTTGAATGATCCCTTAGCATATATCAAGGCTCTTGTCTCTTTTGGATCTTTGG 445
 7292 GTTGAATGATCCCTTATATCAATATGTAATGGGCTTCTGTGCTTTTGGATCTTTGG 7341

466 GTTTAAAGTCGTTTTATCAGAGAGTTGGATTGCAA 501
 7342 GTTTAAAGTCGTTTTATCAGAGACTAGGATTCGAA 7377

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RESULT 4
US-09-764-877-2218
; Sequence 2218, Application US/03764877
; Patent No. US20020147140A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: P0305
; CURRENT APPLICATION NUMBER: US/09/764,877
; CURRENT FILING DATE: 2001-02-17
; Prior application data removed - refer to PAK or file wrapper
; NUMBER OF SEQ. ID NOS: 4631
; SOFTWARE: Patent in Ver. 2.0
; SEQ ID NO 2218
; LENGTH: 9192
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-764-877-2218

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Query Match      86.7%  Score 434.4  DB 10  Length 9192;
Best local Similarity 92.1%  Pired.No. 4.7e-105;
Matches 475; Conservative 0; Mismatches 25; Indels 15; Gaps 11;

Cy 1  GGTTCGGTGAGTTCTTCTTAATCCGAGAGTCTCTAGTTTGAATGCACATGCGGCTGAGAGAC 60
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 6862 GGTTCGGTGAGATCTCTTAATCCGAGAGTCTCTAGTTTGAATGCACATGCGGCTGAGAGAT 6921

Cy 61  AGTTCGTTGTAATCTCTCTCTTTTATCTTGTGGAGAGAGTCTTTAGTTCCAACTATGT 120
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 6922 AGTTCGTTAATATTTCTCTCTTTTACATCTGTTGAGAGAGAGTTTACHTTAAATATGT 6981

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[illegible]

Qy	406	GTTCAAATTGATCCCTTTAGCATTAATGATGGCCTCTCTTCTCTTTCAGACTTGTG	465
Db	7282	GTTGAATTGATCCCTTTAGCATTAATGTAATGGCCCTCTTGTCTCTTTTGACTTGTG	7341
Qy	466	GTTTAAAGTCTGTTTATTATCAGAGAGATTGGATTGCAA	501
Db	73342	GTTTAAAGTCTGTTTATTATCAGAGACTAGATTGCAA	73377

RESULT 5
US-10-092-154-1246
; Sequence 1246, Application US/10092154

Publication No. US20030054375A1
GENERAL INFORMATION
APPLICANT: Roser et al.
TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
FILE REFERENCE: PC009C1
CURRENT APPLICATION NUMBER: US/10/332,154
CURRENT FILING DATE: 2002-03-07
NUMBER OF SEQ. NOS: 2003
Prior Application removed - See File Wrapper or Paim
SOFTWARE: PatentIn Ver 2.0

SEQ ID NO: 1246
LENGTH: 5192
TYPE: DNA
ORGANISM: Homo sapiens
J030730-154-1246

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Query Match      26.7%; Score 434.4; DB 14; Length 9:92;
Best Local Similarity 92.1%; Pred. 4.6e-108;
Matches 475; Conservative 0; Mismatches 26; Indels 15; Gaps 1;

      1  GGTTCGCGAGTTCTCAATCTCGATCTCTAGTTGATTGCACCTGGCGCTCAGAGAC 60
      bb  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
    6862  GGTTCGAGTGAGATCTTAATCTCGAGTTCTAGTTCGATTSCACCTGTGGCTCAGAGAT 6921
      yy  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

    61  AGTTCGTTCTAAATCTCTGTCCTTTTACATTTGCTGAGGAGTGCCTTTAGTCCCAACTATGT 120
      yy  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
    5922  AGTTGTCTAATCTCTGTTCTTTTACATTTGCTGAGGAGAGCTTTACTCCAAAGTATGT 6981
      bb  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

    121  GTCGAATTTTGGAATAGTGTGGTGCTGAGAGAAATGTTATATCTCTTCATCTGG 180
      yy  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
    6982  GGTCAATTTTGGAATAGTGTGGTGCTGAGAGAAATGTTATATCTCTTCATCTGG 7041
      bb  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

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7042	GGTGGAGAGTCTGAGATGCTATTAGCTCGCTGGTGAGAGCTGAGTCAATTCCT	7182	TTCTGCTCTGTTGATCTGCTAAATATTGACAGTGGGGTGTTGA	285	GGATATCCCTGTTAACTTCTCTCGGTGATCTCTCTAAATTGTGACAGTGGGGTGTTAA	7161	AGTCTCCCAAGTATTGTGGGGAGTCTAAGTCTCTCTGTAGGCTCTAGGCACTTGCT	345	ACTCTCCCAATTATTGTGTGGGAGTCTAAGTCTCTCTGTAGGCTCTAAAGCACTTGCT	7221	TTATGAGATCTGGGTCTCTCTGTAATGGGTGGATATATTATAGAGTAGTTAGCTCTCTCT	405	TTATGAAATCTGGTGCTCTGTATGGGTGGATATATTATAGAGTAGTTAGCTCTCTCT	7281	GTGGAATGATGCACTTTAGCATATATGATGGGCTCTCTCTCTCTCTTTTGATCTTTGTTG	465	GTGGAATGATGCACTTTATCATTTATGTAATGGGCTCTCTCTCTCTTTTGATCTTTGTTG	7341	GTTTAAAGTCTGTTTATCAGAGAGTTGGATTGCA	501	GTTTAAAGTCTGTTTATCAGAGAGTAGGATTGCAA	7377
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1  RESULT 6
2  US-03-764-877-3756
3  Sequence 3756, Application US/09764877
4  Patent No. US202014740A1
5  GENERAL INFORMATION:
6  APPLICANT: Rosen et al.
7  TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
8  FILE REFERENCE: PC005
9  CURRENT APPLICATION NUMBER: US/39/764,877
10  CURRENT FILING DATE: 2001-01-17
11  Prior application data removed - refer to PALM or file wrapper
12  NUMBER OF SEQ ID NOS: 4031
13  SOFTWARE: Patent.n Ver. 2.0
14  SEQ ID NO 3756
15  LENGTH: 5676
16  TYPE: DNA

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ORGANISM: Homo sapiens
US-09-764-855-3756

Query Match 86.1%; Score 431.2; DB 10; Length 5676;
Best Local Similarity 91.7%; Pred. No. 2.8e-107;
Matches 473; Conservative 0; Mismatches 28; Indels 15; Gaps 1;
CY 1 GGTGTTGGTGGAGTCTCTAATCCCTGAGTCTTAGTTTATTCGCACTGGGGCTGAGAGAC 60
DB 3323 GGTGTTGGTGGAGTCTCTAATCCCTGAGTCTTAGTTTATTCGCACTGGGGCTGAGAGAC 3382
CY 61 AGTTGTTGTAATTTCTGTTCTTTTACATTTGCTGAGGAGTGGCTTAGTCCCACTATGT 120
DB 3383 AGTTGTTGTAATTTCTGTTCTTTTACATTTGCTGAGGAGTGGCTTAGTCCCACTATGT 3442
CY 121 GGTCAATTTTGAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
DB 3443 GGTCAATTTTGAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 3502
CY 181 GGTGTTAGAGTCTGTAGATGTTATAGTCTGATGCTGATGCTGATGCTGATGCTGATGCTG 240
DB 3503 GGTGTTAGAGTCTGTAGATGTTATAGTCTGATGCTGATGCTGATGCTGATGCTGATGCTG 3562
CY 241 GGA-----TCTGCTGTTGTTGATGCTGATGCTGATGCTGATGCTGATGCTGATGCTG 285
DB 3563 GGTATCTGTTTAACTCTGCTCATGATGCTGCTCATGATGCTGCTCATGATGCTGCTCATG 3622
CY 286 AGTCTCCAGATTTATTTGGTGGAGTCTAGTCTCTTTGATGCTCTTAGGAGTCTGCT 345
DB 3623 AGTCTCCAGATTTATTTGGTGGAGTCTAGTCTCTTTGATGCTCTTAGGAGTCTGCT 3682
CY 346 TTATGAATCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 405
DB 3683 TTATGAATCTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 3742
CY 406 GTTGAATTCACCCCTTTAGCATTTATATGATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 465
DB 3743 GTTGAATTCACCCCTTTAGCATTTATATGATGCTGCTGCTGCTGCTGCTGCTGCTGCTG 3802
CY 466 GTTAAAGTCTGTTTATACAGAGTCTGATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 501
DB 3803 GTTAAAGTCTGTTTATACAGAGTCTGATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 3838
RESULT 7
US-09-764-855-210/c
; Sequence 210, Application US/09764855
; Publication No. US20020119919A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PAL10C1
; CURRENT APPLICATION NUMBER: US/09/764,855
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 334
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 210
; LENGTH: 32199
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-855-210

Query Match 86.1%; Score 431.2; DB 10; Length 32199;
Best Local Similarity 91.7%; Pred. No. 5.9e-107;
Matches 473; Conservative 0; Mismatches 28; Indels 15; Gaps 1;
CY 1 GGTGTTGGTGGAGTCTCTAATCCCTGAGTCTTAGTTTATTCGCACTGGGGCTGAGAGAC 60
DB 7319 GGTGTTGGTGGAGTCTCTAATCCCTGAGTCTTAGTTTATTCGCACTGGGGCTGAGAGAC 7260
CY 61 AGTTGTTGTAATTTCTGTTCTTTTACATTTGCTGAGGAGTGGCTTAGTCCCACTATGT 120

DB 7259 AGTTGTTGTAATTTCTGTTCTTTTACATTTGCTGAGGAGTGGCTTAGTCCCACTATGT 7200
CY 121 GGTCAATTTTGAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
DB 7199 GGTCAATTTTGAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 7140
CY 181 GGTGTTAGAGTCTGTAGATGCTTATTAGTCCACTTGGTGGCAGAGTGGTGGTGGTGGTGG 240
DB 7139 GGTGAGAGTCTGTAGATGCTTATTAGTCCCGTTGGTGGCAGAGTGGTGGTGGTGGTGG 7080
CY 241 GG-----ATCTGCTGTTGTTGATGCTGCTGATGCTGCTGATGCTGCTGATGCTGCTG 285
DB 7079 GGTATCTCTTGTGACTTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 7020
RESULT 8
US-10-072-349-210/c
; Sequence 210, Application US/10072349
; Publication No. US20030254420A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PAL10C1
; CURRENT APPLICATION NUMBER: US/10/072,349
; CURRENT FILING DATE: 2002-02-11
; Prior Application removed - See file Wrapper or Palm
; NUMBER OF SEQ ID NOS: 334
; SOFTWARE: PatentIn Ver. 3.1
; SEQ ID NO 210
; LENGTH: 32199
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-072-349-210
Query Match 86.1%; Score 431.2; DB 14; Length 32199;
Best Local Similarity 91.7%; Pred. No. 5.9e-107;
Matches 473; Conservative 0; Mismatches 28; Indels 15; Gaps 1;
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DB 7319 GGTGTTGGTGGAGTCTCTAATCCCTGAGTCTTAGTTTATTCGCACTGGGGCTGAGAGAC 7260
CY 61 AGTTGTTGTAATTTCTGTTCTTTTACATTTGCTGAGGAGTGGCTTAGTCCCACTATGT 120
DB 7259 AGTTGTTGTAATTTCTGTTCTTTTACATTTGCTGAGGAGTGGCTTAGTCCCACTATGT 7200
CY 121 GGTCAATTTTGAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
DB 7199 GGTCAATTTTGAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 7140
CY 181 GGTGTTAGAGTCTGTAGATGCTTATTAGTCCACTTGGTGGCAGAGTGGTGGTGGTGGTGG 240
DB 7139 GGTGAGAGTCTGTAGATGCTTATTAGTCCCGTTGGTGGCAGAGTGGTGGTGGTGGTGG 7080
CY 241 GG-----ATCTGCTGTTGTTGATGCTGCTGATGCTGCTGATGCTGCTGATGCTGCTG 285
DB 7079 GGTATCTCTTGTGACTTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 7020

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QY 286 AGTCTCCAGATATTATGTTGGGAGGCTTAAGCTCTTTGTATAGTCTCTAGGAGATTCTT 345
Db 7619 AGTCTCCAGATATTATGTTGGGAGGCTTAAGCTCTTTGTATAGTCTCTAGGAGATTCTT 6949
QY 346 TTATGATCTGGGCTGCTCCGATTGGTGCTATATGATATATTTAGGATAGTACTCTTCTT 405
Db 6959 TTATGATCTGGGCTGCTCCGATTGGTGCTATATGATATATTTAGGATAGTACTCTTCTT 6909
QY 406 GTTGAATGATCCCTTTAGGATATATGATGGCTTCTTTGTCTCTTTGTATAGTCTCTT 465
Db 6899 GTTGAATGATCCCTTTAGGATATATGATGGCTTCTTTGTCTCTTTGTATAGTCTCTT 6849
QY 466 GTTAAAGTCTGTTTATACAGAGAGTTGGATTGCAA 501
Db 6839 GTTAAAGTCTGTTTATACAGAGAGTTGGATTGCAA 6804

RESULT 9
US-10-225-810-26/c
; Sequence 26, Application US/0225800
; Publication No. US20030157512A1
; GENERAL INFORMATION:
; APPLICANT: Berrington, Jr., John R.
; TITLE OF INVENTION: Translators and Methods of Using Translators
; FILE REFERENCE: McLaughlin-07:65
; CURRENT APPLICATION NUMBER: US/10/225,810
; CURRENT FILING DATE: 2002-08-21
; NUMBER OF SEQ ID NOS: 76
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 26
; LENGTH: 250000
; TYPES: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (33774)..(433774)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (429533)..(43052)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (45557)..(45656)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (48203)..(48302)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (49552)..(49650)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (51561)..(51660)
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; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (52722)..(52921)
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; NAME/KEY: misc feature
; LOCATION: (53864)..(53963)
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; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (55290)..(55389)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (56674)..(56773)
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; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (57879)..(57978)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (78952)..(79051)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (85316)..(85415)
; OTHER INFORMATION: n is a, c, g, or t
; US-10-225-810-26

Query Match      86.1%   Score 431.2;   DB 12;   Length 250000;
Best Local Similarity 91.7%   Pred. No. 1.4e-106;
Matches 473;   Conservative 0;   Mismatches 28;   Indels 15;   Gaps 2;

QY 1 GGTITTCGGTGGATTTCTTAATCCTGAGTCTCTAGTCTCTAGTCTCTAGTCTCTAGTCTCT 60
Db 124172 GGTITTCGGTGGATTTCTTAATCCTGAGTCTCTAGTCTCTAGTCTCTAGTCTCTAGTCTCT 124113
QY 61 AGTTTGTGTAATTTCTTCTTTTACATTTTGTGAGGAGTCTTTAGTCTCTAGTCTCTAGTCTCT 120
Db 124112 AGTTTGTGTAATTTCTTCTTTTACATTTTGTGAGGAGTCTTTAGTCTCTAGTCTCTAGTCTCT 124053
QY 121 GGTCAATTTTGGAAATAGGTGTGTGTGGTGCTGAGAGAAATGATATTTCTGTTGATTGG 180
Db 124052 GGTCAATTTTGGAAATAGGTGTGTGTGGTGCTGAGAGAAATGATATTTCTGTTGATTGG 123993
QY 181 GGTTTAGAGTCTGTAGATGCTATTAGGTCCTACTTGGTCAGAGCTGGTCCAGAGCTGAGTCTAGTCTCT 240
Db 123992 GGTGGAGAGTCTGTAGATGCTATTAGGTCCTACTTGGTCAGAGCTGGTCCAGAGCTGAGTCTAGTCTCT 123933
QY 241 GGA-----TCTGCTCTTCTGATCTGTCTAAATATGACAGTGGGGTGTGTA 285
Db 123932 GGAATCTCTGTTAACTTCTGCTCTCAATTCATCTGCTCAATGTTGACAGTGGGGTGTGTA 123873
QY 286 AGTCTCCAGTATTATTGTGGGAGTCTAAGTCTCTTTTAGTCTCTAGGAGCTTCT 345
Db 123872 AGTCTCCAGTATTATTGTGGGAGTCTAAGTCTCTTTTAGTCTCTAGGAGCTTCT 123813
QY 346 TTATGAATCTGGGCTGCTCTGATTGGTGCTATATATCTAGGATAGTACTCTTCTT 405
Db 123812 TTATGAATCTGGGCTGCTCTGATTGGTGCTATATATCTAGGATAGTACTCTTCTT 123753
QY 406 GTTGAATGATCCCTTTAGGATATATGATGGCTCTTTTGTCTCTTTTGTATCTTTGTTG 465
Db 123752 GTTGAATGATCCCTTTAGGATATATGATGGCTCTTTTGTCTCTTTTGTATCTTTGTTG 123693
QY 466 GTTAAAGTCTGTTTATACAGAGAGTTGGATTGCAA 501
Db 123692 GTTAAAGTCTGTTTATACAGAGAGTTGGATTGCAA 123657

RESULT 10
US-09-764-801-7392/c
; Sequence 1392, Application US/09764891
; Publication No. US20030077808A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC006
; CURRENT APPLICATION NUMBER: US/09/764,891
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 10231
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 7392
; LENGTH: 5446
; TYPE: DNA
; ORGANISM: Homo sapiens
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US-09-764-891-7392

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Query Match      85.7%; Score 429.6; DB 11; Length 5446;
Best Local Similarity 91.5%; Pred. No. 7.5e-107;
Matches 472; Conservative 0; Mismatches 29; Indels 15; Gaps 1;

QY 1 GGTATTGGAGTTCTTAATCCAGATTCAGTTGATTCGACTGTGGCTGAGAGAC 60
DB 2091 GGTATTGGAGTTCTTAATCCAGATTCAGTTGATTCGACTGTGGCTGAGAGAC 2032
QY 61 AGTTTGTGTAATTCCTGTTTACATTTGCTGAGGAGTCTTATTTCCACTATGT 120
DB 2031 AGTTTGTGTAATTCCTGTTTACATTTGCTGAGGAGTCTTATTTCCACTATGT 1912
QY 121 GGTCAATTTTGAATAGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGG 180
DB 1971 GGTCAATTTTGAATAGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGG 1912
QY 181 GGTATTAGAGTCTGTAGATGCTGATTCAGTCCACTTGTGGTGTGGTGTGGTGTGG 240
DB 1911 GGTGAGAGATTCGTAGATGCTGATTCAGTCCACTTGTGGTGTGGTGTGGTGTGG 1912
QY 241 GG-----ATCTGCTGTTGATGCTGCTGATTCGCTAATATGACAGTGGGTGTGA 285
DB 1851 GGTATCTCTGTTGACTTCTGCTGCTGATTCGCTAATATGACAGTGGGTGTGA 285
QY 286 AGTCTCCAGATTAATTTGTGGAGTCTTAAGTCTCTTTGTAGGTCTCTAGGACTTGT 345
DB 1791 AGTCTCCAGATTAATTTGTGGAGTCTTAAGTCTCTTTGTAGGTCTCTAGGACTTGT 1712
QY 346 TTATGAATCTGGGTGCTGCTGATTCGGTGTGCATATATTTAGGATAGTTAGCTCTTCT 405
DB 1731 TTATGAATCTGGGTGCTGCTGATTCGGTGTGCATATATTTAGGATAGTTAGCTCTTCT 1612
QY 406 GTTGAATTCGATCCCTTAGCATTAATATGATGAGTCTCTTTGCTCTTTTATGATTTTGTG 465
DB 1671 GTTGAATTCGATCCCTTAGCATTAATATGATGAGTCTCTTTGCTCTTTTATGATTTTGTG 1612
QY 466 GTTTAAAGTCTGTTTATCAGAGAGTGGATTCGAA 501
DB 1611 GTTTAAAGTCTGTTTATCAGAGAGTGGATTCGAA 1576

RESULT 11
US-09-764-877-3755
; Sequence 3755, Application US/09764877
; Publication No. US2002014740A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PCC05
; CURRENT APPLICATION NUMBER: US/09/764,877
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - refer to PALM or file wrapper
; NUMBER OF SEQ ID NOS: 4031
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3755
; LENGTH: 5671
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-877-3755
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Query Match      85.7%; Score 429.6; DB 10; Length 5671;
Best Local Similarity 91.5%; Pred. No. 7.6e-107;
Matches 472; Conservative 0; Mismatches 29; Indels 15; Gaps 1;

QY 1 GGTATTGGAGTTCTTAATCCAGATTCAGTTGATTCGACTGTGGCTGAGAGAC 60
DB 3319 GGTATTGGAGTTCTTAATCCAGATTCAGTTGATTCGACTGTGGCTGAGAGAC 3377
QY 61 AGTTTGTGTAATTCCTGTTTACATTTGCTGAGGAGTCTTATTTCCACTATGT 120
DB 3378 AGTTTGTGTAATTCCTGTTTACATTTGCTGAGGAGTCTTATTTCCACTATGT 3437
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QY 121 GGTCAATTTTGGAAATAGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGG 180
DB 3439 GGTCAATTTTGGAAATAGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGG 2497
QY 181 GGTATTAGAGTCTGTAGATGCTTATTTAGTCCACTTGGTGTGGAGGCTGAGTTCTCT 240
DB 3499 GGTGGAGAGTCTGTAGATGCTTATTTAGTCCACTTGGTGTGGAGGCTGAGTTCTCT 3557
QY 241 GGA-----TCTGCTCTTGTGATCTGTCTAAATATGACAGTGGGTGTGA 285
DB 3559 GGTATATCTTGTAACTCTCTCTCATTCATCTGCTAAATATGACAGTGGGTGTGA 3617
QY 286 AGTCTCCAGATTAATTTGTGGAGTCTTAAGTCTCTTTGTAGGTCTCTAGGACTTGT 345
DB 3616 AGTCTCCAGATTAATTTGTGGAGTCTTAAGTCTCTTTGTAGGTCTCTAAGGACTTGT 3677
QY 346 TTATGAATCTGGGTGCTGCTGATTCGGTGTGCATATATTTAGGATAGTTAGCTCTTCT 405
DB 3676 TTATGAATCTGGGTGCTGCTGATTCGGTGTGCATATATTTAGGATAGTTAGCTCTTCT 3737
QY 406 GTTGAATTCGATCCCTTAGCATTAATATGATGAGTCTCTTTGCTCTTTTATGATTTTGTG 465
DB 3738 GTTGAATTCGATCCCTTAGCATTAATATGATGAGTCTCTTTGCTCTTTTATGATTTTGTG 3797
QY 466 GTTTAAAGTCTGTTTATCAGAGAGTGGATTCGAA 501
DB 3798 GTTTAAAGTCTGTTTATCAGAGAGTGGATTCGAA 3833

RESULT 12
US-09-764-891-10041
; Sequence 10041, Application US/09764891
; Publication No. US2003007780A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PCC06
; CURRENT APPLICATION NUMBER: US/09/764,891
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 1023
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 10041
; LENGTH: 20746
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-891-10041
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Query Match      85.7%; Score 429.6; DB 11; Length 20746;
Best Local Similarity 91.5%; Pred. No. 1.3e-106;
Matches 472; Conservative 0; Mismatches 29; Indels 15; Gaps 1;

QY 1 GGTATTGGAGTCTTCTTAATCTGATGCTCTAGTTGATTCGACTGTGGCTGAGAGAC 60
DB 18494 GGTATTGGAGTCTTCTTAATCTGATGCTCTAGTTGATTCGACTGTGGCTGAGAGAC 18553
QY 61 AGTTTGTGTAATTCCTGTTTACATTTGCTGAGGAGTCTTATTTCCACTATGT 120
DB 18554 AGTTTGTGTAATTCCTGTTTACATTTGCTGAGGAGTCTTATTTCCACTATGT 18613
QY 121 GGTCAATTTTGGAAATAGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGG 180
DB 18614 GGTCAATTTTGGAAATAGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGGTGTGG 18673
QY 181 GGTATTAGAGTCTGTAGATGCTTATTTAGTCCACTTGGTGTGGAGGCTGAGTTCTCT 240
DB 18674 GGTGGAGAGTCTGTAGATGCTTATTTAGTCTGGTGTGGTGTGGTGTGGTGTGGTGTGG 18733
QY 241 GGA-----TCTGCTCTTGTGATCTGTCTAATATGACAGTGGGTGTGA 285
DB 18734 GGTATATCTTGTTAATGTTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 18793
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QY 285 AGTCTCCAGATATTATGTTGGAGTCTAAATGCTTTGTTAGTCTCTAGGACTTCTT 345
DB 18794 AGTCTCCAGATATTATGTTGGAGTCTAAATGCTTTGTTAGTCTCTAGGACTTCTT 38853
QY 346 TTATGAATCTGGTCTCTCTGTTATGGGTGCATATATATTAGATAGTTAGTCTCTCTT 405
DB 19954 TTATGAATCTGGTCTCTCTGTTATGGGTGCATATATATTAGATAGTTAGTCTCTCTT 38923
QY 406 GTTGAATTCATCCCTTTAGCATTAATATGATGGCTCTCTTCTCTCTTTTATCTTTCTTG 465
DB 18914 GTTGAATTCATCCCTTTAGCATTAATATGATGGCTCTCTTCTCTCTTTTATCTTTCTTG 18973
QY 466 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 501
DB 18974 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 19029

RESULT 13
US-09-818-512-3
; Sequence 3, Application US/098:5512
; Patent No. US20020142416A1
; GENERAL INFORMATION:
; APPLICANT: BEASLEY, Ellen et al.
; TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC
; TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CLO01:9231V
; CURRENT APPLICATION NUMBER: US/09:818,512
; CURRENT FILING DATE: 2001-03-28
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 116592
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(116592)
; OTHER INFORMATION: n = A,T,C or G
US-09-818-512-3

Query Match: 65.4%; Score 428; DB 10; Length 116592;
Best Local Similarity 91.3%; Pred. No. 7.6e-106;
Matches 471; Conservative 0; Mismatches 30; Indels 15; Gaps 1;

QY 1 GGTCTTGGTGAGTTCTTAATCTGAGTCTGTTTCATTCACATTCGAGAGTGGGCTCAGAGAC 60
DB 51828 GGCCTTGGTGAGATCTTAATCTGAGTCTGTTTCATTCACATTCGAGAGTGGGCTCAGAGAC 51889
QY 61 AGTTTCTGTAAATTTCTGTTCTTTTACATTTGCTGAGAGTSCCTTAGTTCGCAACTATGT 120
DB 51868 AGTTTGTATAATTTCTGTTCTTTTACATTTGCTGAGAGATCTTAGTTCGCAACTATGT 51947
QY 121 GGTCAATTTTGGATAGAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
DB 51948 GGTCAATTTTGGATAGAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 52007
QY 181 GGTTCAGATCTCTGATAGTCTGTTCTTACATTTGCTGAGAGTGGGCTGAGTTCAGTTCCT 240
DB 52008 GGTGGAGATCTCTGATAGTCTGTTCTTACATTTGCTGAGAGTGGGCTGAGTTCAGTTCCT 52067
QY 241 GG-----ATCTGCTCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTG 285
DB 52068 GGTATCTCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTG 52127
QY 286 AGTCTCCAGATATTATGTTGGAGTCTAAATGCTTTGTTAGTCTCTAGGACTTCTT 345
DB 52128 AGTCTCCAGATATTATGTTGGAGTCTAAATGCTTTGTTAGTCTCTAGGACTTCTT 52187
QY 346 TTATGAATCTGGTCTCTCTGTTATGGGTGCATATATATTAGATAGTTAGTCTCTTCTT 405
DB 52188 TTATGAATCTGGTCTCTCTGTTATGGGTGCATATATATTAGATAGTTAGTCTCTTCTT 52247
QY 406 GTTGAATTCATCCCTTTAGCATTAATATGATGGCTCTCTTCTCTCTTTTATCTTTCTTG 465
DB 52248 GTTGAATTCATCCCTTTAGCATTAATATGATGGCTCTCTTCTCTCTTTTATCTTTCTTG 52307
QY 466 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 501
DB 52308 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 52343
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QY 456 GTTGAATTCATCCCTTTAGCATTAATATGATGGCTCTCTTCTCTCTTTTATCTTTCTTG 465
DB 52248 GTTGAATTCATCCCTTTAGCATTAATATGATGGCTCTCTTCTCTCTTTTATCTTTCTTG 52307
QY 466 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 501
DB 52308 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 52343

RESULT 14
US-10-354-065-3
; Sequence 3, Application US/10354065
; Publication No. US20030138837A1
; GENERAL INFORMATION:
; APPLICANT: BEASLEY, Ellen et al.
; TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC
; TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CLO01:9231V
; CURRENT APPLICATION NUMBER: US/10:354,065
; CURRENT FILING DATE: 2003-01-30
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 116592
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(116592)
; OTHER INFORMATION: n = A,T,C or G
US-10-354-065-3

Query Match: 85.4%; Score 428; DB 12; Length 116592;
Best Local Similarity 91.3%; Pred. No. 7.6e-106;
Matches 471; Conservative 0; Mismatches 30; Indels 15; Gaps 1;

QY 1 GGTCTTGGTGAGTTCTTAATCTGAGTCTGTTTCATTCACATTCGAGAGTGGGCTCAGAGAC 60
DB 51828 GGCCTTGGTGAGATCTTAATCTGAGTCTGTTTCATTCACATTCGAGAGTGGGCTCAGAGAC 51887
QY 61 AGTTTCTGTAAATTTCTGTTCTTTTACATTTGCTGAGAGTSCCTTAGTTCGCAACTATGT 120
DB 51868 AGTTTGTATAATTTCTGTTCTTTTACATTTGCTGAGAGATCTTAGTTCGCAACTATGT 51947
QY 121 GGTCAATTTTGGATAGAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
DB 51948 GGTCAATTTTGGATAGAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 52007
QY 181 GGTTCAGATCTCTGATAGTCTGTTCTTACATTTGCTGAGAGTGGGCTGAGTTCAGTTCCT 240
DB 52008 GGTGGAGATCTCTGATAGTCTGTTCTTACATTTGCTGAGAGTGGGCTGAGTTCAGTTCCT 52067
QY 241 GG-----ATCTGCTCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTG 285
DB 52068 GGTATCTCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTGATCTGTTG 52127
QY 286 AGTCTCCAGATATTATGTTGGAGTCTAAATGCTTTGTTAGTCTCTAGGACTTCTT 345
DB 52128 AGTCTCCAGATATTATGTTGGAGTCTAAATGCTTTGTTAGTCTCTAGGACTTCTT 52187
QY 346 TTATGAATCTGGTCTCTCTGTTATGGGTGCATATATATTAGATAGTTAGTCTCTTCTT 405
DB 52188 TTATGAATCTGGTCTCTCTGTTATGGGTGCATATATATTAGATAGTTAGTCTCTTCTT 52247
QY 406 GTTGAATTCATCCCTTTAGCATTAATATGATGGCTCTCTTCTCTCTTTTATCTTTCTTG 465
DB 52248 GTTGAATTCATCCCTTTAGCATTAATATGATGGCTCTCTTCTCTCTTTTATCTTTCTTG 52307
QY 466 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 501
DB 52308 GTTAAAGTCTCTTTTATCAGAGACTTGGATTGCAA 52343
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RESULT 15
US10-027-983-11/c
; Sequence 11, Application US10027983
; Publication No. US2003013936CA1
; GENERAL INFORMATION:
; APPLICANT: Kenneth W. Doble
; TITLE OF INVENTION: ANTISENSE MODULATION OF ESTROGEN RECEPTOR ALPHA EXPRESSION
; FILE REFERENCE: R/S-0340
; CURRENT APPLICATION NUMBER: US10/027,983
; CURRENT FILING DATE: 2001-12-18
; NUMBER OF SEQ ID NOS: 99
; SEQ ID NO 11
; LENGTH: 392000
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: unsure
; LOCATION: 13742
; OTHER INFORMATION: unknown
; NAME/KEY: unsure
; LOCATION: 13742
; OTHER INFORMATION: unknown
; NAME/KEY: misc feature
; LOCATION: 138122...138221
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: unsure
; LOCATION: 145507
; OTHER INFORMATION: unknown
; NAME/KEY: unsure
; LOCATION: 151967
; OTHER INFORMATION: unknown
; NAME/KEY: misc feature
; LOCATION: 151967...1542066
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: unsure
; LOCATION: 154217
; OTHER INFORMATION: unknown
; NAME/KEY: misc feature
; LOCATION: 154337...154336
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 174571...174756
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 186224...186323
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 195242...195341
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: unsure
; LOCATION: 202703
; OTHER INFORMATION: unknown
; NAME/KEY: misc feature
; LOCATION: 202771...202870
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 205246...2125602
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 216126...218225
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 220363...220459
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 222717...222816
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 223391...224080
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
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; LOCATION: (227487)...227586;
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: (230157)...230256;
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: (232299)...232398;
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: (236552)...236651;
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: (238769)...248788;
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: exon
; LOCATION: (118288)...119101;
; OTHER INFORMATION: exon 1C
; NAME/KEY: exon:intron junction
; LOCATION: (151129)...151130;
; OTHER INFORMATION: exon 5:intron 5
; NAME/KEY: exon:intron junction
; LOCATION: (299249)...299249;
; OTHER INFORMATION: exon 9:intron 9
; NAME/KEY: exon:intron junction
; LOCATION: (348578)...348579;
; OTHER INFORMATION: exon 10:intron 10
; NAME/KEY: intron
; LOCATION: (348579)...381838;
; OTHER INFORMATION: intron 1C
; NAME/KEY: intron:exon junction
; LOCATION: (396185)...396186;
; OTHER INFORMATION: intron 11:exon 12
[US10-027-983-11]

Query Match      85.4%   Score 428; DB 12; Length 392000;
Best Local Similarity 91.3%   Pred. No. 1.3e-105;
Matches 471; Conservative 3; Mismatches 30; Indels 15; Gaps 1;

QY      1  GGTTCGCGTGAGTTCTTAAATCTGAGTCTCTAGTTGATGGACCTGTGSCCTGAGAGAC 60
DB      318449  GGTTCGAGTGAGTTCTTAAATCTGAGTCTCTAGTTGATGGACCTGTGSCCTGAGAGAT 318390
QY      61  AGTTTGTTGTAATTCCTGTTCTTTTACATTTGCTGAGGAGTGCTTTAGTTCCCAACTATGT 120
DB      318359  AGTTTGTTGTAATTCCTGTTCTTTTACATTTGCTGAGGAGTGCTTTAGTTCCCAACTATGT 318320
QY      121  GGTCATTTTGGAAATAGGTGCTGGTGTGCTGCTGAGAGAAATGTATATTCCTGTTGATTGG 180
DB      318329  GGTCATTTTGGAAATAGGTGCTGGTGTGCTGAGAGAAATGTATATTCCTGTTGATTGG 318270
QY      181  GGTTCAGATTCGTAGATGCTTATTAGTCCACTTGTGTCAGAGCTGAGTTCAGTTCCT 240
DB      318269  GGTTCAGATTCGTAGATGCTTATTAGTCCACTTGTGTCAGAGCTGAGTTCAGTTCCT 318210
QY      241  GGA-----TCTGTCCTTTGTCATCTGCTAATATTGACAGTGGGTGTGA 285
DB      318209  GGATATCCTTGTTAATTTCTGTCCTGTCATCTGCTAATATTGACAGTGGAGTGTAA 318150
QY      286  AGTCCCAAGTATTATGTTGGAGTCTAAGTCTCTTTTAGGTCTCTAGGACATTGCT 345
DB      318149  AGTCCCAATTAATTGTTGGAGTCTAAGTCTCTTTTAGGTCTCTAAGGACATTGCT 318090
QY      346  TTATGAATCTGGTGCTCTGTAATGGTGCAATATATATTAGGATAGTAGTCTCTCT 405
DB      318089  TTATGAATCTGGTGCTCTGTAATGGTGCAATATATATTAGGATAGTAGTCTCTCT 318030
QY      406  GTTGAATTGATCCCTTAGCATTATATGATGGCTCTTTGCTCTTTTGTGATCTTTGTC 465
DB      318029  GTTGAATTGATCCCTTAGCATTATATGATGGCTCTTTGCTCTTTTGTGATCTTTGTC 317970
QY      466  GTTTAAAGTCTGTTTATCAGAGATTGGATTGCAA 501
DB      317969  GTTTAAAGTCTGTTTAAACAGAGACTAGGATTGCAA 317934
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Job time : 130.415 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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(without alignments)
11633.329 Million cell updates/sec

Title: US-09-830-902-1_COPY_50000_50500

Perfect score: 501
Sequence: 1 ggttttgcgagttcttta.....atcagagagtgatggcan 501

Scoring table: IDENTITY NJC
Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0
Maximum DB seq length: 200000000
Post-processing: Minimum Match 3%
Maximum Match 100%
Listing first 45 summaries

Database: EST:

- 1: em_estba.*
- 2: em_esthum.*
- 3: em_estlin.*
- 4: em_estmu.*
- 5: em_estov.*
- 6: em_estp1.*
- 7: em_estro.*
- 8: em_hic.*
- 9: gb_est1.*
- 10: gb_est2.*
- 11: gb_hic.*
- 12: gb_est3.*
- 13: gb_est4.*
- 14: gb_est5.*
- 15: em_estfur.*
- 16: em_estom.*
- 17: em_gss_hur.*
- 18: em_gss_inv.*
- 19: em_gss_p.h.*
- 20: em_gss_vri.*
- 21: em_gss_fvr.*
- 22: em_gss_nam.*
- 23: em_gss_mus.*
- 24: em_gss_pro.*
- 25: em_gss_rtd.*
- 26: em_gss_pmg.*
- 27: em_gss_vrl.*
- 28: gb_gss1.*
- 29: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
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C 2	426.4	85.1	1075	11	BC012129	Homo sapi
C 3	424.8	84.8	880	10	BC619398	602084576
C 4	423.8	84.6	2495	11	BC037952	Homo sapi

C 5	423.2	84.5	609	2	HS074999	
C 6	422.4	84.3	579	3	AL13263	
C 7	422.4	84.3	579	3	AL13377	
C 8	421.2	84.1	651	28	AC385776	
C 9	420.6	84.0	778	13	B0547312	
C 10	420.6	83.8	1100	12	BM473445	
C 11	420.6	83.8	3314	28	AF101615	
C 12	418.4	83.5	729	29	B272880	
C 13	415.4	82.9	604	12	B018378	
C 14	415.4	82.9	609	12	B1021312	
C 15	415.4	82.9	626	12	B1019813	
C 16	415.2	82.3	750	9	AL223802	
C 17	414.2	82.7	701	2	HS037915	
C 18	413.6	82.6	777	14	CB312020	
C 19	412.8	82.4	688	10	EG572005	
C 20	412.8	82.2	1037	29	B2601843	
C 21	410.4	81.9	609	10	AW860981	
C 22	410.4	81.9	829	29	B2774229	
C 23	410.4	81.8	712	28	AQ055327	
C 24	406.4	81.1	799	14	CD511598	
C 25	404.8	80.8	651	29	AG019019	
C 26	404.8	80.8	904	9	AV711080	
C 27	402.2	80.3	698	10	BE173627	
C 28	399.6	79.8	574	2	HS073550	
C 29	399.6	79.8	707	28	AQ470254	
C 30	398.8	79.6	497	28	AQ428674	
C 31	395.5	79.0	687	10	BE173465	
C 32	395	78.8	554	10	BF676389	
C 33	393.4	78.5	829	10	BG537539	
C 34	392.6	78.4	615	10	AW856417	
C 35	392.6	78.4	654	9	AV730783	
C 36	392.2	78.3	658	29	AG034925	
C 37	392.2	78.3	732	29	BZ612233	
C 38	390	77.8	709	13	EC84643	
C 39	389.6	77.8	663	29	AG119932	
C 40	388.2	77.5	668	29	AQ422387	
C 41	388.2	77.5	701	29	AG116167	
C 42	385	76.8	657	29	AG019827	
C 43	384.6	76.8	703	28	BZ917429	
C 44	382	76.2	623	28	AQ555093	
C 45	381.8	75.2	909	29	BZ598445	

ALIGNMENTS

RESULT 1
BU959549/c
LOCUS
DEFINITION
IMAGE: 6737882 5'.. mRNA sequence.
AGENCOURT_0623586 NIH MGC_127 Homo sapiens cDNA clone
753 bp -RNA linear EST 21-OCT-2002
BU959549: G:24189121
AGENCOURT
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-tr@mail.nih.gov
Tissue Procurement: NCI
CDNA Library Preparation: Michael Brownstein Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: JLCM368 row: k column: 01
High quality sequence stop: 526.

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FEATURES
  source
    Location/Qualifiers
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        /organism="Homo sapiens"
        /mol_type="rRNA"
        /db_xref="taxon:9606"
        /clone="IMAGE:6737882"
        /tissue_type="mixed (pool of 40 RNAs)"
        /lab_hosts="DH10B (T4-phage-resistant)"
        /clone_lib="NIH_MGC_127"
        /notes="Vector: pNR-LIB; Site 1: Sfil (ggccattatggcc);
          Site 2: Sfil (ggccgcctggcc); Double-stranded cDNA was
          prepared from a pool of 40 cell line polyA+ RNAs (Bladder
          - 2%, blood - 33.4%, brain - 5.4%, breast - 12.5%, colon -
          4%, connective tissue - 1.4%, eye - 1%, intestine - 2.5%,
          kidney - 2.2%, liver - 5.7%, lung - 20.8%, NK-cell - 5.2%,
          ovary - 4%, pharynx - 2.5%, prostate - 4.3%, salivary
          gland - 1.3%, and skin - 2.3%). 5' and 3' adaptors were
          used in cloning as follows:
          5'-AAGCAGTGGTATCAGCCAGAGTGGCCATTAGCGCGG-3' and
          5'-ATTCTAGAGCGGCGGCGCCGACATG-3'. Full-length
          enriched library was constructed using the Clontech
          Creator SMART kit and size-selected to contain the 1-2 kb
          size fraction (other fractions present in NIH_MGC_127 and
          NIH_MGC_128). Library created in the laboratory of T.
          Usdin, M.D., Ph.D. (NIH). Note: this is a NIH_MGC
          Library."
      305 a 170 c 130 q 146 t 2 others
      5'..1075
        /organism="Homo sapiens"
        /mol_type="rRNA"
        /clone="IMAGE:4575709"
        /tissue_type="Primary B-Cells from Tonsils"
        /clone_lib="NIH_MGC_48"
        /lab_hosts="DH10B-R"
        /notes="Vector: pOTB7"

BASE COUNT
ORIGIN
  Query Match 95.1%; Score 426.4; DB 13; Length 753;
  Best Local Similarity 91.1%; Pred. No. 4.7e-64;
  Mismatches 47%; Conservative 3; Indels 15; Gaps 1;

  QY 1 GGTTCGCGAGTCTCTTAAAGCTGAGTCTTACCTTATGTTGAGTCTGAGAGAC 60
  DB 571 GGTTCGAGTGGTTCCTTAAAGCTGAGTCTTACCTTATGTTGAGTCTGAGAGAC 512
  QY 61 AGTTCTGTTAAATCTGTTCTTTACATTTCTGAGAGAGTCTTATGTTGAGTCTG 120
  DB 511 AGTTCTGTTAAATCTGTTCTTTACATTTCTGAGAGAGTCTTATGTTGAGTCTG 452
  QY 121 GGTCAATTTTGGAAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180
  DB 451 GGTCAATTTTGGAAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 392
  QY 181 GGTTCAGAGTCTGTTAGAGTCTTATGTTGAGTCTGTTGAGTCTGTTGAGTCTG 240
  DB 391 GGTTCAGAGTCTGTTAGAGTCTTATGTTGAGTCTGTTGAGTCTGTTGAGTCTG 332
  QY 241 GGA-----TCTCTCTGTTGATCTGCTAATATGACAGTGGGGTGTGA 285
  DB 331 GGTATCTCTGTTAACTGCTGCTCACTGATCTGACTAATGTTGACAGTGGGTGAA 272
  QY 286 AGTCTCCAGATATTATGTTGGGAGTCTAAGTCTCTTTGAGGTTCTAGGAGTCTGT 345
  DB 271 AGTCTCCAGATATTATGTTGGGAGTCTAAGTCTCTTTGAGGTTCTAAGGAGTCT 212
  QY 345 TATGATGAGTGGGTGCTGCTGATATGGTGGTCAATATATTTAGGATAGTATCTCT 405
  DB 211 TATGATGAGTGGGTGCTGCTGATATGGTGGTCAATATATTTAGGATAGTATCTCT 152
  QY 406 GTGAATTCATCCCTTAGCATATATGATGCTCTCTCTCTCTCTCTCTCTCTCTCT 465
  DB 151 GTTGATTCATCCCTTAGCATATATGATGCTCTCTCTCTCTCTCTCTCTCTCTCT 92
  QY 466 GTTAAAGTCTGTTTATCAGAGAGTGGATTGCA 501
  DB 91 GTTAAAGTCTGTTTATCAGAGAGTGGATTGCA 56

  RESULT 2
  BC012129/c 1075 bp rRNA 1:near HTC 04-MAR-2003
  LOCUS

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DEFINITION Homo sapiens, Similar to LINE retrotransposable element 1, clone IMAGE:4575709, mRNA.

ACCESSION BC012129

VERSION BC012129.1 GI:15082424

KEYWORDS HTC.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE

AUTHORS Strausberg R.

TITLE Direct Submission

JOURNAL Submitted (02-AUG-2001) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2500, USA

REMARK

CONVENT NIH-MGC Project URL: <http://mgc.nci.nih.gov>

Contact: MGC help desk

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: Louis Staudt

cDNA Library Preparation: Rubin Laboratory

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (ILNL)

cDNA Sequencing by: Genome Sequence Centre,

BC Cancer Agency, Vancouver, BC, Canada

RefSeqAccession: U00000

Steven Jones, Jennifer Asano, Ian Bosdet, Yaron Butterfield, Susanna Chan, Feachan Chiu, Chris Pfeil, Erin Gaffan, Ron Guin, Patricia Hsiang, Martin Krzywinski, Reta Kutsche, Oliver Lee, Soo Sen Lee, Victor Ling, Carrie Mathewson, Cardice McLeavy, Steven Ness, Pawan Pandoh, Anra-Liisa Prabhur, Parvaneh Saeedi, Jacqueline Schein, Duane Smalhus, Michael Smith, Joraine Spence, Jeff Scott, Michael Thorne, Miranada Tsai, Kandasja van den Bosch, Jill Vardy, George Yang, Scott Zuyerdun, Marco Maria.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/ILNL at: <http://image.llnl.gov>

Series: IPAL Plate: 29 Row: 5 Column: 10

This clone was selected for full length sequencing because it passed the following selection criteria: Similarity but not identity to protein

This clone has the following problem: retained intron.

Location/Qualifiers

1..1075

/organism="Homo sapiens"

/mol_type="rRNA"

/db_xref="taxon:9606"

/clone="IMAGE:4575709"

/tissue_type="Primary B-Cells from Tonsils"

/clone_lib="NIH_MGC_48"

/lab_hosts="DH10B-R"

/notes="Vector: pOTB7"

BASE COUNT 451 a 239 c 179 g 206 t

ORIGIN

Query Match 85.1%; Score 426.4; DB 11; Length 1075;

Best Local Similarity 91.1%; Pred. No. 4.5e-64;

Mismatches 47%; Conservative 0; Mismatches 31; Indels 15; Gaps 1;

QY 1 GGTTCGCGAGTCTCTTAAAGCTGAGTCTTACCTTATGTTGAGTCTGAGAGAC 60

DB 835 GGTTCGCGAGTCTCTTAAAGCTGAGTCTTACCTTATGTTGAGTCTGAGAGAT 776

QY 61 AGTTGTTTAAATCTGTTCTTTACATTTCTGAGGAGTCTTATGTTGAGTCTG 120

DB 775 AGTTGTTTAAATCTGTTCTTTACATTTCTGAGGAGTCTTATGTTGAGTCTG 716

QY 121 GGTCAATTTTGGAAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 180

DB 715 GGTCAATTTTGGAAATAGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 656

QY 181 GGTTCAGAGTCTGTTAGAGTCTTATGTTGAGTCTGTTGAGTCTGTTGAGTCTG 240

DB 655 GGTTCAGAGTCTGTTAGAGTCTTATGTTGAGTCTGTTGAGTCTGTTGAGTCTG 596


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Db      146 GTGGAATGACCTTACCATATGTAAGGCTCTTGTGCTATTGATCTTGGTGTG 87
QY      466 GTTAAAGTCTGTTTATCAGACAGATGATTGCAA 501
Db      86 GTTAAAGTCTGTTTATCAGACAGATGATTGCAA 51

RESULT 6
LOCUS   AL119263/c
DEFINITION DKF2p76:K0913 r1 761 (synonym: hamy2) Homo sapiens cDNA clone
ACCESSION AL119263
VERSION   AL119263.1 GI:5925162
KEYWORDS  EST.
SOURCE    Homo sapiens (human)
ORGANISM  Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 579)
AUTHORS   Ansoerge,W., Benes,V., Krieger,S., Mewes,H.W., Gassenhuber,J. and
           Wiemann,S.
TITLE      EST (Ansoerge, Benes, et al.)
JOURNAL    Unpublished
COMMENT    Contact: Ansoerge W
           MIPS
           Ingolstaedter Landstr.1, D-85764 Neuherberg, Germany
           This is the 5' sequence of the clone insert
           Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
           Research Center (DKFZ); Email: s.wiemann@dkfz-heidelberg.de;
           sequenced by EMBL within the cDNA sequencing consortium of the
           German Genome Project.
           No 5' sequence available.
           This clone is available at the RZPD in Berlin.
           Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
           Berlin-Charlottenburg, GERMANY; Email: c.cone@rpd.de.
           Location/Qualifiers
             loc=579
             /organism="Homo sapiens"
             /mol_type="mRNA"
             /db_xref="taxon:9606"
             /clone="DKF2p76:K0913"
             /tissue_type="amygdala"
             /dev_stage="adult"
             /lab_host="DH10B"
             /clone_lib="761" (synonym: hamy2)
             /notes="Vector: pSport1; Site_1: Not1; Site_2: SalI"
BASE COUNT 243 a 136 c 91 g 109 t
ORIGIN
Query Match      84.3% Score 422.4; DB 9; Length 579;
Best Local Similarity 91.0%; Pred. No. 2.4e-63;
Matches 466; Conservative 0; Mismatches 31; Indels 15; Gaps 1;

QY      5 TTGCGTGAAGTCTTCTTAATCTCTGAGTCTCTAGTTTGAATTGCACCTGTGGCCTGAGACAGATT 64
Db      579 TTGAGTGAAGTCTTCTTAATCTCTGAGTCTCTAGTTTGAATTGCACCTGTGGCCTGAGACAGATT 520

QY      65 TGTTGTAAATCTCTGTTCTTTTACATTTGCTGAGGAGTGGTTAGTTCGCAACTATGTGGTC 124
Db      519 TGTTGAATTTCTGTTCTTTTACATTTGCTGAGGAGTGGTTAGTTCGCAACTATGTGGTC 460

Query Match      84.3% Score 422.4; DB 9; Length 579;
Best Local Similarity 91.0%; Pred. No. 2.4e-63;
Matches 466; Conservative 0; Mismatches 31; Indels 15; Gaps 1;

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Db      339 ATGCTGTAAATCTCTGTTCTCTCATTTGATCTGTCTAATGTGACAGTGGGGTGTGAAGTC 280
QY      290 TCCAGTATTATTGCTGGGAGTCTAAAGTCTCTTTTGTAGGCTCTAGGACATTGCTTTAT 349
Db      279 TCCAGTATTATTGCTGGGAGTCTAAAGTCTCTTTGTAGGCTCTAGGACCTGCTTTAT 220
QY      350 GAATCTGGGTGCTCTGTTATGGTGCATATATATTAGGATAGTATGCTCTTCTTGTG 409
Db      219 GAATCTGGGTGCTCTGTTATGGTGCATATATATTAGGATAGTATGCTCTTCTTATG 160
QY      410 AATGTATCTCTTACGATATATGATGGCTCTTGTCTCTTTTGATCTTTTGTGGATT 469
Db      259 AATCTATCTCTTACGATATATGATGGCTCTTGTCTCTTTTGATCTTTTGTGGATT 100
QY      470 AAAGTCTGTTTATCAGACAGATGATTGCAA 501
Db      99 AAAGTCTGTTTATCAGACAGATGATTGCAA 68

RESULT 7
LOCUS   AL133777/c
DEFINITION DKF2p76:K12113 r1 761 (synonym: hamy2) Homo sapiens cDNA clone
ACCESSION AL133777
VERSION   AL133777.1 GI:6601965
KEYWORDS  EST.
SOURCE    Homo sapiens (human)
ORGANISM  Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 579)
AUTHORS   Ansoerge,W., Winkler,J., Mewes,H.W., Weil,B. and Wiemann,S.
TITLE      EST (Ansoerge,W., Winkler,J., Mewes,H.W., Weil,B. and Wiemann,S.)
JOURNAL    Unpublished
COMMENT    Contact: Ansoerge W
           MIPS
           Ingolstaedter Landstr.1, D-85764 Neuherberg, Germany
           This is the 5' sequence of the clone insert
           Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
           Research Center (DKFZ); Email: s.wiemann@dkfz-heidelberg.de;
           sequenced by ENBL (European Molecular Biology Laboratories,
           Heidelberg/Germany); within the cDNA sequencing consortium of the
           German Genome Project.
           No 5' sequence available.
           This clone (DKF2p76:K12113) is available at the RZPD in Berlin.
           Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
           Berlin-Charlottenburg, GERMANY; Email: c.cone@rpd.de.
           Location/Qualifiers
             loc=579
             /organism="Homo sapiens"
             /mol_type="mRNA"
             /db_xref="taxon:9606"
             /clone="DKF2p76:K12113"
             /tissue_type="amygdala"
             /dev_stage="adult"
             /lab_host="DH10B"
             /clone_lib="761" (synonym: hamy2)
             /notes="Vector: pSport1; Site_1: Not1; Site_2: SalI"
BASE COUNT 243 a 136 c 91 g 109 t
ORIGIN
Query Match      84.3% Score 422.4; DB 9; Length 579;
Best Local Similarity 91.0%; Pred. No. 2.4e-63;
Matches 466; Conservative 0; Mismatches 31; Indels 15; Gaps 1;

QY      5 TTGCGTGAAGTCTTCTTAATCTCTGAGTCTCTAGTTTGAATTGCACCTGTGGCCTGAGACAGATT 64
Db      579 TTGAGTGAAGTCTTCTTAATCTCTGAGTCTCTAGTTTGAATTGCACCTGTGGCCTGAGACAGATT 520

QY      65 TGTTGTAAATCTCTGTTCTTTTACATTTGCTGAGGAGTGGTTAGTTCGCAACTATGTGGTC 124
Db      519 TGTTGAATTTCTGTTCTTTTACATTTGCTGAGGAGTGGTTAGTTCGCAACTATGTGGTC 460

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High quality sequence step: 613.
Location/Qualifiers
1..778

FEATURES
source

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5613573"
/lab_host="DH10B (11 phage-resistant)"
/clone_lib="NIH MGC 82"
/notes="Organ: testis; Vector: pCMV-SB (Clontech); Site: 1: Sfil (ggcgctggcc); Site 2: Sfil (ggcattatggcc); 3' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CACGCCAATATGGCC-3' and 3' adaptor sequence: 5'-ATTCTAGAGCCGAGCGCGCCGACATG-dT(30)BN-3' (where B = A, C, or G and N = A, C, or T). Average insert size 1.35 kb (range 0.9-4.0 kb). 14/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA)."
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BASE COUNT 326 a 173 c 129 g 146 t 4 others
ORIGIN

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Query Match: 84.0%; Score 420.6; DB 13; Length 778;
Best Local Similarity 90.5%; Pred. No. 4.6e-63;
Matches 466; Conservative 0; Mismatches 34; Indels 15; Gaps 1;

QY 2 GTTTGGTGAGTCTCTTAATCTGAGTCTCTAGTTGATTCACATGCTGGCCCTGAGAGACA 61
DB 650 GTTTGAGTGAGTCTCTTAATCTGAGTCTCTAATTTGATTGCACTGGCTGCTGATAGATA 591
QY 62 GTTTGTTGTAATCTCTCTTTACATTTGCTGAGGAGTGGTTAGTTCCAACTATGTG 121
DB 590 GTTTGTTGTAATCTCTCTTTACATTTGCTGAGGAGGCTTACTTCCAACTATGTG 531
QY 122 GTCAATTTGGAAATAGTGTGTGTGCTGCTGAGAGAAATGATATCTGTTGATTTGGG 181
DB 530 GTCAATTTGGAAATAGTGTGTGTGCTGCTGAGAGAAATGATATCTGTTGATTTGGG 471
QY 182 GTTACAGTCTGACATGCTAATTAAGTCCACTGGCTGACAGGTGATTCAGTCTGCTG 241
DB 470 GTGAGAGTCTGATAGTGTCTAATTAAGTCTCTTTGGTGCAGAGGTGATTCAGTCTG 411
QY 242 GA-----CTGTCTCTTTGATCTGTCTAATAATGACAGTGGGCTTGAA 286
DB 410 GATATCTTTGTTAACTTATGCTCTTTGATCTGTCTAATGCTGACAGTGGGCTTGAA 351
QY 287 GTCTCCAGTATTAATGTGTGGAGTCTAAGTCTCTTTAGGTCTCTTACAGACTTGCTT 346
DB 350 GTCTCCAGTATTAATGTGTGGAGTCTAAGTCTCTTTAGGTCTCTTACAGACTTGCTT 291
QY 347 TATGAATCTGGGTGCTCTGTATGGGTGCATATATATTAGATAGTATAGCTTCTCTG 406
DB 290 TATGAATCTGGGTGCTCTGTATGGGTGCATATATATTAGATAGTATAGCTTCTCTG 231
QY 407 TTGAATGATCCCTTATAGATATATAGATAGGCTCTCTCTCTTTGATCTCTTTGATCTT 466
DB 230 TTGAATGATCCCTTATAGATATATAGATAGGCTCTCTCTCTTTGATCTCTTTGATCTT 171
QY 467 TTTAAACTCTGTTTATATCAGAGAGTTGGATTGCAA 501
DB 170 TTTAAAGTCTGTTTATATCAGAGTCTAGGATTGCAA 136
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RESULT 10
BM473445/c
LOCUS BM473445 1100 bp mRNA linear EST 05-FEB-2002
DEFINITION AGENCOURT_6484467 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:5613573
5', mRNA sequence.
ACCESSION BM473445
VERSION BM473445.1 GI:118522487
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 bases 1 to 1100
NIH-MGC http://imgc.ncl.nih.gov/
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strussberg, Ph.D.
E-mail: rstruss@fmail.nih.gov
Tissue Procurement: ATCC/DCTP/STP
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNU)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM1223C row: a column: 24
High quality sequence step: 716.
Location/Qualifiers
1..1100
/organism="Homo sapiens"
/mcl_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5613573"
/tissue_type="melanotic melanoma"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC 72"
/notes="Organ: skin; Vector: pCMV-SORT6; Site 1: NotI;
Site 2: SalI; cloned unidirectionally. Primer: Oligo dt.
Average insert size 2 kb. Library constructed by Life
Technologies."

BASE COUNT 467 a 246 c 177 g 209 t 1 others
ORIGIN

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Query Match: 83.8%; Score 420; DB 12; Length 1100;
Best Local Similarity 90.3%; Pred. No. 5.6e-63;
Matches 466; Conservative 0; Mismatches 35; Indels 15; Gaps 1;

QY 1 GGTTTGCGTGAATCTCTTAATCTGAGTCTCTAGTTGATTCACATGCTGGCTGGAGAGAC 50
DB 685 GGTTTGAGTGAGTCTCTTAATCTGAGTCTCTAGTTGATTCACATGCTGGCTGGAGAT 626
QY 61 AGTTTGTGTAAATTTCTGTCTCTTTACATTTGCTGAGGAGTCTTTAGTTCCAACTATGT 120
DB 625 AGTTTGTATAATCTCTGTATTTACATTTGCTGAGGAGAGCTTACTTCCAACTATGT 566
QY 121 GGTCAATTTGGAAATAGTGTGTGTGCTGCTGAGAGAAATGATATCTCTGTGATTTGG 180
DB 565 GGTCAATTTGGAAATAGTGTGTGTGCTGCTGAGAGAAATGATATCTCTGTGATTTGG 506
QY 181 GGTTTAGAGTCTGTGATGATCTTATAGGTCCACTTGGTGCAGAGCTGAGTTCACTTCT 240
DB 505 GGTGGAGAGTCTGTGATGATCTTATAGGTCTCTTGGTGCAGAGCTGAGTTCACTTCT 446
QY 241 GG-----ATCTGTCTTTGTTGATCTGTCTAATAATTTGACAGTGGGCTTGAA 285
DB 445 GGTATAGTCTTTGTTGATCTGTCTTATAGTCTGTCTAATAATTTGACAGTGGGCTTGAA 366
QY 286 AGTCTCCAGTATATAGTGTGGAGTCTAAGTCTCTTTTGTAGTCTCTAGGACTTCT 345
DB 365 AGTCTCCAGTATATAGTGTGGAGTCTAAGTCTCTTTTGTAGTCTCTAGGACTTCT 326
QY 346 TATGAATCTGGGTGCTCTGTATTTGGGTGCATATATATTAGGATAGTATAGCTTCTCT 405
DB 325 TATGAATCTGGGTGCTCTGTATTTGGGTGCATATATATTAGGATAGTATAGCTTCTCT 266
QY 406 GTTGAATTTGATCCCTTTAGCATATATAGTGGCTCTCTTTGCTCTTTTGTATCTTTGTTG 465
DB 265 GTTGAATTTGATCCCTTTAGCATATATAGTGGCTCTCTTTGCTCTTTTGTATCTTTGTTG 206
QY 466 GTTTAAAGTCTGTTTATATCAGAGAGTTGGATTGCAA 501
DB 205 GTTTAAAGTCTGTTTATATCAGAGACTAGGATTGCAA 170
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RESULT 11
AF101615/c
LOCUS      AF101615 Human Homo sapiens genomic clone pTWS233.20, genomic
DEFINITION survey sequence.
ACCESSION  AF101615
VERSION    AF101615.1 GI:424977.8
KEYWORDS   GSS.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 3324)
AUTHORS   Bepko G., O'Brian, K.C., Kim, Y.C., Schreiber, G. and Pitterle, D.M.
TITLE     A 1.4-Mb high-resolution physical map and contig of chromosome
            segment 1p15.5 and genes in the LCR1A metastasis suppressor
            region
JOURNAL   Genomics 55 (2): 164-175 (1999)
MEDLINE   9914294
PUBMED   9933563
COMMENT   Contact: Bepko G.
            Duke University Medical Center
            Box 2616, MSRB, Room 117, Durham, NC 27710, USA
            part of a 1.4 megabase contig including the LCR1A metastasis
            suppressor region Bin 8
            Class: unknown.
            Location/Qualifiers
                1..3314
                /organism="Homo sapiens"
                /mol_type="genomic DNA"
                /db_xref="taxon:9606"
                /map="1p15.5"
                /clone="pTWS233.20"
                /clone_lib="Human"
BASE COUNT  1300 a 743 c 672 g 601 t
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Query Match      83.8%; Score 420; DB 20; Length 3314;
Best Local Similarity 90.3%; Pred. No. 4,9e-63;
Matches 466; Conservative 0; Mismatches 35; Indels 15; Gaps 1;
QY      1  GGTTCGGTGGAGTTCCTTAACTCCAGTTCAGTTGATTGACGTGGCTGAGAGAC 60
DB      1  GGTTCGGTGGAGTTCCTTAACTCCAGTTCAGTTGATTGACGTGGCTGAGAGAC 2559
QY      61  AGTTTCGTGTAATTCCTGTTCTTTTACATTTCTGAGGAGTCTTTAGTCCCAACTATGT 120
DB      2558  AGTTTCGTGTAATTCCTGTTCTTTTACATTTCTGAGGAGTCTTTAGTCCCAACTATGT 2499
QY      121  GGTCAATTTTGGAAATAGGTGTGGTGGTGGCTGAGGAGATGATATTCCTGATTGG 180
DB      2498  GGTCAATTTTGGAAATAGGTGTGGTGGTGGCTGAGGAGATGATATTCCTGATTGG 2439
QY      181  GGTTCAGAGTCTGTAGATGCTCTATTAGGCTCCATTTGGTGGAGAGCTGAGTTCAGTTCCT 240
DB      2439  GGTGGAGAGTCTGTAGATGCTCTATTAGGCTCCATTTGGTGGAGAGCTGAGTTCAGTTCCT 2379
QY      241  GSA-----CTGTCCTTTGTTGATCTGTCTAATATTGACAGTGGGGTGTGA 285
DB      2378  GGATACTCTTGTGAATTTCTGCTGCTGATCTGTCTAATATTGACAGTGGGGTGTGA 2319
QY      286  AGTCTCCAGTATATTGTGTGGAGCTCAAGTCTCTTTGTAGGCTCTTAGGAGCTTCT 345
DB      2318  AGTCTCCAGTATATTGTGTGGAGCTCAAGTCTCTTTGTAGGCTCTTAGGAGCTTCT 2259
QY      346  TTATGAATCTGGGTGCTCTGATTCGGGTGCATATATATTAGGATAGTACGCTCTCT 405
DB      2258  TTATGAATCTGGGTGCTCTGATTCGGGTGCATATATATTAGTATAGTATGTTCTCT 2199
QY      406  GTTGAATCTGATCCCTTTAGCAATATATGATGGGCTCTTTGCTCTCTTTTATCTTTGTTG 465
DB      2199  GTTGAATCTGATCCCTTTAGCAATATATGATGGGCTCTTTGCTCTCTTTTATCTTTGTTG 465

D3      2138  CTTGATATAGATCCCTTTGCGCATATATGATGAGGCTTTCTTTGATCTCTTTGATCTCTTTG 2139
QY      466  GTTTAAAGTCTGTTTATATCAGAGAGTGGATTGCAA 501
DB      2138  GTTTAAAGTCTGTTTATATCAGAGACTAGGATTGCAA 2103

RESULT 12
BZ772880
LOCUS      BZ772880 mcv56c10.b1 HFOSM2007 Homo sapiens genomic, genomic survey
DEFINITION sequence.
ACCESSION  BZ772880
VERSION    BZ772880.1 GI:28946564
KEYWORDS   GSS.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 729)
AUTHORS   Cook, J., Delehaunty, K., Fewell, G., Fulton, L., Megrini, V., Vardis, E.,
            Miner, T., Nash, N., Williams, D. and Wilson, R.K.
TITLE     Homo sapiens Fosmid End Reads
COMMENT   Contact: Richard K. Wilson
            Genome Sequencing Center
            Washington University School of Medicine
            Email: submissions@watson.wustl.edu
            Plate: mcv56 row: c column: 10
            Class: fosmid ends
            High quality sequence start: 38
            High quality sequence stop: 550.
            Location/Qualifiers
                1..729
                /organism="Homo sapiens"
                /mol_type="genomic DNA"
                /db_xref="taxon:9606"
                /clone_lib="HFOSMID307"
                /note="Vector: pccitfos; Site: 1: Eco72i; Human whole
            genome fosmid library was prepared at Washington
            University Genome Sequencing Center. DNA was sheared for
            blunt-ended ligation into pccitfos inducible vector. DNA
            was ordered from Coriell Cell Repository's DNA
            polymorphism discovery resource."
BASE COUNT  145 a 120 c 172 g 289 t
ORIGIN
Query Match      83.5%; Score 418.4; DB 29; Length 729;
Best Local Similarity 90.1%; Pred. No. 1.1e-62;
Matches 465; Conservative 0; Mismatches 36; Indels 15; Gaps 1;
QY      1  GGTTCGGTGGAGTTCCTTAACTCCAGTTCAGTTGATTGACGTGGCTGAGAGAC 60
DB      204  GGTTCGGTGGAGTTCCTTAACTCCAGTTCAGTTGATTGACGTGGCTGAGAGAT 263
QY      61  AGTTTCGTGTAATTCCTGTTCTTTTACATTTGCTGAGGAGTGTTCCTTAACTTCT 120
DB      264  AGTTTCGTGTAATTCCTGTTCTTTTACATTTGCTGAGGAGTGTTCCTTAACTTCT 323
QY      121  GGTCAATTTTGGAAATAGGTGTGGTGGTGGCTGAGGAGATGATATTCCTGATTGG 180
DB      324  GGTCAATTTTGGAAATAGGTGTGGTGGTGGCTGAGGAGATGATATTCCTGATTGG 383
QY      181  GGTTCAGAGTCTGTAGATGCTCTATTAGGCTCCACTTGGTGCAGAGCTGAGTTCAGTTCCT 240
DB      384  GGTGCAGAGTCTGTAGATGCTCTAATAGGTCCTGCTGGTGGTGCAGAGCTGAGTTCAGTTCCT 443
QY      241  GG-----ATCTGCTCTGCTGATCTGTCTAATATTGACAGTGGGGTGTGA 285
DB      444  GGTATCTCTGCTGATCTCTGCTGCTGATCTGTCTAATATTGACAGTGGGGTGTGA 503
QY      286  AGTCTCCAGTATATTGTGTGGAGTCTAAGTCTCTTTGATGCTCTTTGATGAGGACTTCT 345
DB      503  AGTCTCCAGTATATTGTGTGGAGTCTAAGTCTCTTTGATGCTCTTTGATGAGGACTTCT 345
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Db 504 AGTCTCCCATTAATATGTGGGAGCTAAGCTCTCTTTAGTCTCACTCAGGACTCGCT 563
 Cy 346 TTATGAACTGGGTGCGCCCTGATCGGTGCAATATATTAGCATAGTATAGTCTCTCT 428
 Db 564 TTATGAACTGGGTGCGCCCTGATCGGTGCAATATATTAGCATAGTATAGTCTCTCT 428
 Cy 406 GTTGAATGATCCCTTTAGCATATATGATGGCCCTCTCTCTCTCTCTCTCTCTCTCT 465
 Db 624 GTTGAATGATCCCTTTAGCATATATGATGGCCCTCTCTCTCTCTCTCTCTCTCTCT 465
 Cy 466 GTTGAATGATCCCTTTAGCATATATGATGGCCCTCTCTCTCTCTCTCTCTCTCTCT 501
 Db 684 GTTGAATGATCCCTTTAGCATATATGATGGCCCTCTCTCTCTCTCTCTCTCTCTCT 519

RESULT 13
 LOCUS B1018878/c 604 bp mRNA linear EST 14-JUN-2001
 DEFINITION IL3-MT0267-150101-435-B02 MT0267 Homo sapiens cDNA, mRNA sequence.
 ACCESSION B1018878
 VERSION B1018878.1 GI:14425508
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE
 AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 604).
 Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Bata,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare ,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.
 Shotgun sequencing of the human transcriptome with ORF expressed sequence tags

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000); MEDLINE 20202663
 PUBMED 10737800
 COMMENT Contact: Simpson A.J.G.
 Laboratory of Cancer Genetics
 Ludwig Institute for Cancer Research
 Rua Prof. Antonio Prudente 109, 4 andar, 01506-010, Sao Paulo-SP, Brazil
 Tel.: +55-11-2704322
 Fax: +55-11-2707001
 Email: asimpson@ludwig.org.br
 This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
 (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=IL3&st=IL3-MT0267-050101-435-B02&3=2001-01-05&4=1)
 Seq primer: puc 18 forward
 High quality sequence stop: 603.
 Location/Qualifiers
 ..604
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /dev_stage="Adult"
 /catalytic="MT0267"
 /note="Organ: marrow; Vector: puc18; Site: 1; Site 2: Site 3: A mini-library was made by cloning products derived from ORESTES PCR (J.S. Letters Patent application No. 196 /716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

BASE COUNT 253 a 142 c 92 g 117 t
 ORIGIN
 Query Match 82.9%; Score 425.4; DA 12; Length 604;
 Best Local Similarity 90.9%; Pred. No. 3.8e-62;
 Matches 470; Conservative C; Mismatches 31; Indels 16; Gaps 2;

Cy 1 GGTTTGGTGAGTCTCTTAATCCCTGAGTCTTAGTTGATTCGACTCGGCTGAGAGAC 60
 Db 596 GGTTTGAGTGGATCTTAATCCCTGAGTCTTAGTTGATTCGACTCGGCTGAGAGAT 537
 Cy 61 AGTTTGTGTAAATTTCTCTCTTTTACATTGGCTGAGGAGTGCCTTTAGTCCAAATATGT 120
 Db 536 AGTTTGTGTAAATTTCTCTCTTTTACATTGGCTGAGGAGACTTTATTTCCAAAGTATGT 477
 Cy 121 GTCATTTTGGATAGTGTGGTGTGGTGTG-AGAAGAAATGATATTTCTTTGATTTG 179
 Db 478 GTCATTTTGGATAGTGTGGTGTGGTGTG-GAAGAAATGATATTTCTTTGATTTG 417
 Cy 180 GGGTTTAGAGTCTCTGATAGTGTATTAGTCCAGCTGGTGCAGAGCTGAGTCAATTC 239
 Db 418 GGGTGGAGAGTCTCTGATAGTGTATTAGTCCAGCTGGTGCAGAGCTGAGTCAATTC 357
 Cy 240 TGG-----ATCTGCTTTTGTATCTGCTTAATATTGACAGTGGGGTGTG 284
 Db 358 TGGGTATCTGTTGTAACCTTCTGCTCTGATCTGCTAATGTTGACATTTGGGGTGTGA 297
 Cy 295 AAGTCTCCAGTATATTTGTGTGGAGTCTAAGTCTTTTGTAGGTCTCTAGGAGACTTGC 344
 Db 296 AAGTCTCCAGTATATTAATGTGTGGAGTCTAAGTCTTTTGTAGGTCTCTAGGAGACTTGC 237
 Cy 345 TTATGAATCTGGTGGTCTCTGATTGGTGATATATATTAGGATAGTCTAGTCTCTCT 404
 Db 236 TTATGAATCTGGTGGTCTCTGATTGGTGATATATATTAGGATAGTCTAGTCTCTCT 177
 Cy 405 TTTGAATGATCCCTTTTAGCATATATGATGGCTCTCTTTGTCTTTGATCTCTTTGTT 464
 Db 176 TTTGAATGATCCCTTTTAGCATATATGATGGCTCTCTTTGTCTTTGATCTCTTTGTT 117
 Cy 465 GGTTTAAAGTCTGTTTATACAGAGAGTGGATTGGAA 501
 Db 116 GGTTTAAAGTCTGTTTATACAGAGAGTGGATTGGAA 80

RESULT 14
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 DEFINITION IL3-MT0267-150101-448-C10 MT0267 Homo sapiens cDNA, mRNA sequence.
 ACCESSION B1021312
 VERSION B1021312.1 GI:14427942
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 609).
 Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Bata,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare ,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.
 Shotgun sequencing of the human transcriptome with ORF expressed sequence tags

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000); MEDLINE 20202663
 PUBMED 10737800
 COMMENT Contact: Simpson A.J.G.
 Laboratory of Cancer Genetics
 Ludwig Institute for Cancer Research
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
 Tel.: +55-11-2704322
 Fax: +55-11-2707001
 Email: asimpson@ludwig.org.br
 This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
 (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=IL3&st=IL3-MT0267-150101-448-C10&3=2001-01-15&4=1)
 Seq primer: puc 18 forward

TITLE
 JOURNAL
 MEDLINE
 PUBMED
 COMMENT
 BASE COUNT 253 a 142 c 92 g 117 t
 ORIGIN
 Query Match 82.9%; Score 425.4; DA 12; Length 604;
 Best Local Similarity 90.9%; Pred. No. 3.8e-62;
 Matches 470; Conservative C; Mismatches 31; Indels 16; Gaps 2;

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High quality sequence stop: 609.
FEATURES
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      /organism="Homo sapiens"
      /mol_type="mRNA"
      /db_xref="taxon:9606"
      /clone_lib="WT0267"
      /note="Organ: marrow; Vector: puc18; Site 1: SmaI; Site 2: SmaI; A mini-library was made by cloning products derived from ORFESTS PCR (U.S. letters Patent application No. 196 716 - Ludwig Institute for Cancer Research) profiles into the puc18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."
BASE COUNT      255 a      143 c      93 g      118 t
ORIGIN
Query Match      82.9%; Score 415.4; DB 12; Length 629;
Best Local Similarity 90.9%; Pred. No. 3.7e-62;
Matches 470; Conservative 0; Mismatches 31; Indels 16; Gaps 2;
CY 1 GGTCTTGGCTGAGTTCTTCAATCCGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGAC 60
DB 596 GGTCTTGGCTGAGTCTTCAATCCGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGAT 537
CY 61 AGTTCTTGGCTGAGTCTTCAATCCGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGAT 120
DB 596 AGTTCTTGGCTGAGTCTTCAATCCGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGAT 537
CY 536 AGTTCTTGGCTGAGTCTTCAATCCGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGAT 427
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CY 121 GGTCAATTTGGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGATGATGTTGATGTTG 173
DB 476 GGTCAATTTGGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGATGATGTTGATGTTG 417
CY 180 GGTCTTGGCTGAGTCTTCAATCCGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGAT 239
DB 416 GGTCTTGGCTGAGTCTTCAATCCGAGTGGTGTGATGTTGATGCTGAGTGGCTGAGAGAT 357
CY 240 TGG-----ATCTGCTGTTGATGCTGCTGATGCTGATGCTGATGCTGATGCTGATGCTG 284
DB 356 TGGTATCTGCTGATGCTGATGCTGATGCTGATGCTGATGCTGATGCTGATGCTGATGCTG 297
CY 285 AGCTCTCCAGTATTATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 344
DB 296 AGCTCTCCAGTATTATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 237
CY 345 TTATGAATCTGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 404
DB 236 TTATGAATCTGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 177
CY 405 TGTGTAATGATCCCTTTACCATATATATGATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 464
DB 176 TGTGTAATGATCCCTTTACCATATATATGATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 117
CY 465 GGTCTTAAAGTCTGTTTATCAGAGAGTGGATGCA 501
DB 116 GGTCTTAAAGTCTGTTTATCAGAGAGTGGATGCA 80
RESULT 15
BIC19813
LOCUS      BIC19813      625 bp      mRNA      linear      EST 14-JUN-2001
DEFINITION IL3-WT0267-110101-442-F12 MT0267 Homo sapiens cDNA, mRNA sequence.
ACCESSION  BIC19813
VERSION    BIC19813.1 GI:14426443
KEYWORDS  EST.
SOURCE    Homo sapiens (human);
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 626)
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Brunes,M.R.,
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Cosra,F.F.,
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